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FEATURE: unsure
NAME/KEY: unsure
LOCATION: (653)
OTHER INFORMATION: Unidentified at time of filing

FEATURE: unsure
NAME/KEY: unsure
LOCATION: (656)
OTHER INFORMATION: Unidentified at time of filing

FEATURE: unsure
NAME/KEY: unsure
LOCATION: (660)
OTHER INFORMATION: Unidentified at time of filing

FEATURE: unsure
NAME/KEY: unsure
LOCATION: (661)
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FEATURE: unsure
NAME/KEY: unsure
LOCATION: (662)
OTHER INFORMATION: Unidentified at time of filing

Query Match 93.1%; Score 27; DB 3; Length 888;
Best Local Similarity 83.3%; Pct. No. 1.3e+03;
Matches 5; Conservative 1; Mismatches 0; Indels 8

US-09-351-215-4

Qy      1 SSGFSL 6
Db     248 SSGFSL 253

RESULT 13
US-08-658-857B-16
Sequence 16, Application US/08658857B
Patent No. 6040435
GENERAL INFORMATION:
APPLICANT: Hancock, Robert E. W.
APPLICANT: Karunaratne, Nedra
TITLE OF INVENTION: ANTIMICROBIAL CATIONIC PEPTIDES
NUMBER OF SEQUENCES: 37
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson, P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: BASESEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/658, 857B
FILING DATE: May 31, 1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/460, 464
FILING DATE: June 2, 1995
ATTORNEY/AGENT INFORMATION:
NAME: Halle, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE DOCUMENT NUMBER: 07420/014001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5099
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 20 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-658-857B-16

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RESULT 14
US-08-763-226C-16
; Sequence 16, Application US/08763226C
; Patent No. 6057291
; GENERAL INFORMATION:
;    APPLICANT: Hancock, Robert E. W.
;    APPLICANT: Karunaratne, Nedra
;    TITLE OF INVENTION: ANTIMICROBIAL CATIONIC PEPTIDES
;    NUMBER OF SEQUENCES: 39
;    CORRESPONDENCE ADDRESS:
;    ADDRESSEE: Fish & Richardson P.C.
;    STREET: 4225 Executive Square, Suite 1400
;    CITY: La Jolla
;    STATE: CA
;    COUNTRY: USA
;    ZIP: 92037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/763,226C
; FILING DATE: 10-DEC-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/6558, 857
; FILING DATE: 31-MAY-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Haile, Lisa A.
; REGISTRATION NUMBER: 38,347
; REFERENCE DOCKET NUMBER: 07420/014001
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619/678-5070
; TELEFAX: 619/678-5099
; INFORMATION FOR SEQ ID NO: 16:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 20 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide

US-08-763-226C-16
Query Match          89.7%; Score 26; DB 3; Length 20;
Best Local Similarity 83.3%; Pred. No. 46;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy      1 SSGPSSL 6
Db      12 SSGPSSL 17

RESULT 15
US-09-307-200-16
; Sequence 16, Application US/09307200
; Patent No. 6297215
; GENERAL INFORMATION:
;    APPLICANT: Hancock, Robert E. W.
;    APPLICANT: Karunaratne, Nedra
;    TITLE OF INVENTION: ANTIMICROBIAL CATIONIC PEPTIDES
;    NUMBER OF SEQUENCES: 39
;    CORRESPONDENCE ADDRESS:
;    ADDRESSEE: Fish & Richardson P.C.
;    STREET: 4225 Executive Square, Suite 1400

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STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
OPERATING SYSTEM: Windows 95
SOFTWARE: FASTSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/307,200
FILING DATE:
PRIORITY NUMBER: 08/763,226
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 3B,347
REFERENCE/DOCKET NUMBER: 07420/014001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5070
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 20 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-307-200-16

Query Match 89.7%; Score 26; DB 3; Length 20;
Best Local Similarity 83.3%; Pred. No. 46;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
Qy 1 SSGPST 6
Db 12 SSGPAL 17

Search completed: March 10, 2004, 09:28:53
Job time : 3.32296 secs

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OM protein - protein search, using SW model

Run on: March 10, 2004, 09:16:59 ; Search time 26.7237 Seconds
(without alignments)

268.645 Million cell updates/sec

Title: US-09-848-834A-10
Perfect score: 186
Sequence: 1 FMNFTVSEWLVRPKVSASHLEGPSLHWSYGLRPX 34

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Gapop 10.0 , Gapext 0.5

Searched: 809742 seqs, 211153259 residues

Total number of hits satisfying chosen parameters:

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Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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16: /cgn2_6_ptodata/2/pubpaa/us10_NEWPUB.pep:/*
17: /cgn2_6_ptodata/2/pubpaa/us60_NEWPUB.pep:/*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match	Length	DB ID	Description
1	185	99.5	34	9	US-09-848-834A-10	Sequence 10, Appl
2	185	99.5	50	9	US-09-848-834A-18	Sequence 18, Appl
3	119	64.0	194	14	US-10-295-074-46	Sequence 46, Appl
4	118	63.4	158	14	US-10-297-942-12	Sequence 12, Appl
5	116	62.4	285	14	US-10-295-074-11	Sequence 11, Appl
6	116	62.4	287	14	US-10-295-074-13	Sequence 13, Appl
7	114	61.3	158	14	US-10-297-942-10	Sequence 10, Appl
8	112	60.2	21	9	US-09-943-548-3	Sequence 3, Appl
9	112	60.2	21	9	US-09-848-834A-4	Sequence 4, Appl
10	112	60.2	21	9	US-09-785-215-6	Sequence 6, Appl
11	112	60.2	21	10	US-09-405-986-2	Sequence 2, Appl
12	112	60.2	21	14	US-10-204-362-6	Sequence 3, Appl
13	112	60.2	21	14	US-10-339-522-3	Sequence 8, Appl
14	112	60.2	21	14	US-10-223-711-8	Sequence 6, Appl
15	60.2				US-10-223-809A-6	

RESULT 1
US-09-848-834A-10
; Sequence 10, Application US/09848834A
; Patent No. US2002007616A1
; GENERAL INFORMATION:
/ APPLICANT: Aptagen Corporation
/ TITLE OF INVENTION: Chimeric Peptide Immunogens
/ FILE REFERENCE: 11020165-0007
/ CURRENT APPLICATION NUMBER: US/09-846,834A
/ CURRENT FILING DATE: 2001-05-04
/ PRIOR APPLICATION NUMBER: 60/1202,328
/ PRIORITY FILING DATE: 2000-05-05
/ NUMBER OF SEQ ID NOS: 20
/ SOFTWARE: Patentin version 3.0
/ SEQ ID NO: 10
/ LENGTH: 34
/ TYPE: PRT
/ ORGANISM: Artificial Sequence
/ FEATURE:
/ OTHER INFORMATION: ChimERIC peptide consisting of amino acid sequence 947-967 of t
/ OTHER INFORMATION: Tetanus toxoid precursor (TetoxoLySIN) linked by a spacer to a
/ OTHER INFORMATION: into acid sequence 2-10 of the Tetanus Toxoid Precursor
/ NAME/KEY: MOD-RES
/ LOCATION: (1)..(1)
/ OTHER INFORMATION: Amidated phenylalanine
/ NAME/KEY: PEPTIDE
/ LOCATION: (1)..(21)
/ OTHER INFORMATION: Amino acids 947-967 of the Tetanus Toxoid Precursor
/ NAME/KEY: PEPTIDE
/ LOCATION: (22)..(25)
/ OTHER INFORMATION: Spacer peptide
/ NAME/KEY: PEPTIDE
/ LOCATION: (26)..(34)
/ OTHER INFORMATION: Amino acids 2-10 of the human GnRH hormone
/ NAME/KEY: MOD-RES
/ LOCATION: (34)..(34)
/ OTHER INFORMATION: Amidated glycine or glycynamide

US-09-848-834A-10

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Best Local Similarity 100.0%; Pred. No. 7.5e-18;
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 FNNFTVFWLVPKVSSASHLEGPSLHWSYGRP 33
Db 1 FNNFTVFWLVPKVSSASHLEGPSLHWSYGRP 33

RESULT 2
US-09-848-834A-18

/ Sequence 18, Application US/0948834A
/ GENERAL INFORMATION:
/ APPLICANT: Abtton Corporation
/ TITLE OF INVENTION: Chimeric Peptide Immunogens
/ FILE REFERENCE: 1102865-0047
/ CURRENT APPLICATION NUMBER: US/09/848,834A
/ CURRENT FILING DATE: 2001-05-04
/ PRIOR APPLICATION NUMBER: 60/202,328
/ PRIOR FILING DATE: 2000-05-05
/ NUMBER OF SEQ ID NOS: 20
/ SOFTWARE: Patentin version 3.0
/ SEQ ID NO: 18
/ LENGTH: 50
/ TYPE: PRT
/ ORGANISM: Artificial Sequence
/ FEATURE:
/ OTHER INFORMATION: Chimeric peptide consisting of amino acid sequence 1-10 of human GnRH linked by a spacer to amino acid sequence 947-967 of the Tetanus Toxoid precursor (Tetoxylysin) protein linked by a spacer to other information: o amino acid sequence 2-10 of human GnRH
/ NAME/KEY: MOD_RES
/ LOCATION: (1) ..(1)
/ OTHER INFORMATION: Pyroglutamic acid or 5-oxoproline
/ NAME/KEY: MOD_RES
/ LOCATION: (50) ..(50)
/ OTHER INFORMATION: Amidated glycine or glycaminide
/ NAME/KEY: PEPTIDE
/ LOCATION: (1) ..(10)
/ OTHER INFORMATION: Amino acid sequence 1-10 of the human GnRH hormone
/ NAME/KEY: PEPTIDE
/ LOCATION: (11) ..(16)
/ OTHER INFORMATION: Spacer peptide
/ NAME/KEY: PEPTIDE
/ LOCATION: (17) ..(37)
/ OTHER INFORMATION: Amino acid sequence 947-967 of the Tetanus toxoid precursor (Tetoxylysin)
/ NAME/KEY: PEPTIDE
/ LOCATION: (38) ..(41)
/ OTHER INFORMATION: Spacer peptide
/ NAME/KEY: PEPTIDE
/ LOCATION: (42) ..(50)
/ OTHER INFORMATION: Amino acid sequence 2-10 of the human GnRH hormone
US-09-848-834A-18

Query Match 99.5%; Score 185; DB 9; Length 50;
Best Local Similarity 100.0%; Pred. No. 1.1e-17;
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 FNNFTVFWLVPKVSSASHLEGPSLHWSYGRP 33
Db 17 FNNFTVFWLVPKVSSASHLEGPSLHWSYGRP 49

RESULT 3
US-10-295-074-46

/ Sequence 46, Application US/10295074
/ Publication No. US20030185845A1
/ GENERAL INFORMATION:
/ APPLICANT: Pharmexa A/S

Query Match 63.4%; Score 118; DB 14; Length 158;
Best Local Similarity 100.0%; Pred. No. 4e-08;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 FNNFTVFWLVPKVSSASHLEG 22
Db 133 FNNFTVFWLVPKVSSASHLEG 154

RESULT 5
US-10-295-074-11

/ Sequence 11, Application US/10295074
/ Publication No. US20030185845A1
/ GENERAL INFORMATION:
/ APPLICANT: Pharmexa A/S

Publication No. US20030185845A1
; GENERAL INFORMATION:
; APPLICANT: Pharmexa A/S
; TITLE OF INVENTION: NOVEL IMMUNOGENIC MINIMICS OF MULTIMER PROTEINS
; FILE REFERENCE: P1013DK00
; CURRENT APPLICATION NUMBER: US/10/295,074
; CURRENT FILING DATE: 2002-11-15
; NUMBER OF SEQ ID NOS: 60
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 11
; LENGTH: 285
; TYPE: PRT
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: 2 human IIL5 monomers joined by P2 and P30 epitopes
US-10-295-074-11

Query Match 62.4%; Score 116; DB 14; Length 285;
Best Local Similarity 91.7%; Pred. No. 1.4e-07;
Matches 22; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 FNNFTVSPWLRPKVSKASHLEGPS 24
Db 150 FNNFTVSPWLRPKVSKASHLEPT 173 /

RESULT 6
US-10-295-074-13
; Sequence 13, Application US/10295074
; Publication No. US20030185845A1
; GENERAL INFORMATION:
; APPLICANT: Pharmexa A/S
; TITLE OF INVENTION: NOVEL IMMUNOGENIC MINIMICS OF MULTIMER PROTEINS
; FILE REFERENCE: P1013DK00
; CURRENT APPLICATION NUMBER: US/10/295,074
; CURRENT FILING DATE: 2002-11-15
; NUMBER OF SEQ ID NOS: 60
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 13
; LENGTH: 287
; TYPE: PRT
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Two human IIL5 monomers joined by diglycine linker and including terminal positioned P30 and P2 epitopes
US-10-295-074-13

Query Match 62.4%; Score 116; DB 14; Length 287;
Best Local Similarity 91.7%; Pred. No. 1.4e-07;
Matches 22; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 FNNFTVSPWLRPKVSKASHLEGPS 24
Db 24 FNNFTVSPWLRPKVSKASHLEPT 47

RESULT 7
US-10-297-942-10
; Sequence 10, Application US/10297942
; Publication No. US20030185816A1
; GENERAL INFORMATION:
; APPLICANT: Perking BY
; TITLE OF INVENTION: Solubilised Protein Vaccines
; FILE REFERENCE: P68445US0
; CURRENT APPLICATION NUMBER: US/10/297,942
; CURRENT FILING DATE: 2003-04-21
; PRIOR APPLICATION NUMBER: PCT/DK01/00431
; PRIOR FILING DATE: 2001-10-16
; PRIOR APPLICATION NUMBER: DK PA 2000 00966
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 10

US-09-848-834A-4
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 Best Local Similarity 100.0%; Pred. No. 3e-08; Gaps 0;
 Matches 21; Conservative 0; Mismatches 0; Indels 0;
 SEQ ID NO: 1 FNNFTVSEWLRLVPKVASHLE 21
 ORGANISM: Clostridium tetani
 LENGTH: 21
 SEQ ID NO: 2 1 FNNFTVSEWLRLVPKVASHLE 21
 RESULT 10
 US-09-785-215-6
 Sequence 6, Application US/09785215
 Publication No. US20020187157A1
 GENERAL INFORMATION:
 APPLICANT: JENSEN, Martin Roland et al.
 TITLE OF INVENTION: NOVEL METHOD FOR DOWN-REGULATION OF AMYLOID
 FILE REFERENCE: 3631-0107P
 CURRENT APPLICATION NUMBER: US/09/785,215
 CURRENT FILING DATE: 2001-02-20
 NUMBER OF SEQ ID NOS: 19
 SEQ ID NO: 6
 NUMBER OF SEQ ID NOS: 19
 SEQ ID NO: 6
 LENGTH: 21
 TYPE: PRT
 ORGANISM: Clostridium tetani
 US-09-785-215-6
 Query Match Score 112; DB 9; Length 21;
 Best Local Similarity 100.0%; Pred. No. 3e-08; Gaps 0;
 Matches 21; Conservative 0; Mismatches 0; Indels 0;
 SEQ ID NO: 1 FNNFTVSEWLRLVPKVASHLE 21
 SEQ ID NO: 2 1 FNNFTVSEWLRLVPKVASHLE 21
 RESULT 11
 US-09-405-986-2
 Sequence 2, Application US/09405986
 Publication No. US2003015715A1
 GENERAL INFORMATION:
 APPLICANT: BAY, Sylvie
 APPLICANT: CANTACUZENE, Danièle
 APPLICANT: LECLERC, Claude
 APPLICANT: LO MAN, Richard
 TITLE OF INVENTION: COMPRISING THE SAME AND USE THEREOF
 FILE REFERENCE: 1341 US 35655
 CURRENT APPLICATION NUMBER: US/09/405,986
 CURRENT FILING DATE: 1999-09-27
 EARLIER APPLICATION NUMBER: 60/041,726
 EARLIER FILING DATE: 1997-03-27
 NUMBER OF SEQ ID NOS: 3
 SEQ ID NO: 2
 LENGTH: 21
 TYPE: PRT
 ORGANISM: Clostridium tetani
 US-09-405-986-2
 Query Match Score 112; DB 10; Length 21;
 Best Local Similarity 100.0%; Pred. No. 3e-08; Gaps 0;
 Matches 21; Conservative 0; Mismatches 0; Indels 0;
 SEQ ID NO: 1 FNNFTVSEWLRLVPKVASHLE 21
 SEQ ID NO: 2 1 FNNFTVSEWLRLVPKVASHLE 21
 RESULT 12
 US-10-204-362-6
 Sequence 6, Application US/10204362
 GENERAL INFORMATION:
 APPLICANT: M&B Biotech A/S
 TITLE OF INVENTION: NO. US20030086538A1 Method For Down-Regulation Of Amyloid
 FILE REFERENCE: 3631-0120P
 CURRENT APPLICATION NUMBER: US/10/204,362
 CURRENT FILING DATE: 2002-08-16
 NUMBER OF SEQ ID NOS: 16
 SOFTWARE: PatentIn Ver. 3.0
 SEQ ID NO: 6
 LENGTH: 21
 TYPE: PRT
 ORGANISM: Clostridium tetani
 US-10-204-362-6
 Query Match Score 112; DB 14; Length 21;
 Best Local Similarity 100.0%; Pred. No. 3e-08; Gaps 0;
 Matches 21; Conservative 0; Mismatches 0; Indels 0;
 SEQ ID NO: 1 FNNFTVSEWLRLVPKVASHLE 21
 SEQ ID NO: 2 1 FNNFTVSEWLRLVPKVASHLE 21
 RESULT 13
 US-10-339-522-3
 Sequence 3, Application US/10339522
 Publication No. US20030108559A1
 GENERAL INFORMATION:
 APPLICANT: Bittershaus, Charles W.
 APPLICANT: Thomas, Lawrence J.
 TITLE OF INVENTION: MODULATION OF CHOLESTERYL ESTER TRANSFER PROTEIN (CETP) ACTIVITY
 FILE REFERENCE: TCS-411 LP US-3
 CURRENT APPLICATION NUMBER: US/10/339,522
 CURRENT FILING DATE: 2003-01-08
 PRIOR APPLICATION NUMBER: 08/432,483
 PRIOR FILING DATE: 1995-05-01
 PRIOR APPLICATION NUMBER: PCT/US96/06147
 PRIOR FILING DATE: 1996-05-01
 PRIOR APPLICATION NUMBER: 08/945,289
 PRIOR FILING DATE: 1997-10-17
 PRIOR APPLICATION NUMBER: 09/943,334
 PRIOR FILING DATE: 2001-08-30
 PRIOR APPLICATION NUMBER: 09/943,548
 PRIOR FILING DATE: 2001-08-30
 NUMBER OF SEQ ID NOS: 9
 SEQ ID NO: 3
 LENGTH: 21
 TYPE: PRT
 ORGANISM: Artificial Sequence
 OTHER INFORMATION: helper T cell epitope of tetanus toxin
 US-10-339-522-3
 Query Match Score 112; DB 14; Length 21;
 Best Local Similarity 100.0%; Pred. No. 3e-08; Gaps 0;
 Matches 21; Conservative 0; Mismatches 0; Indels 0;
 SEQ ID NO: 1 FNNFTVSEWLRLVPKVASHLE 21
 SEQ ID NO: 2 1 FNNFTVSEWLRLVPKVASHLE 21
 RESULT 14
 US-10-223-711-8
 Sequence 8, Application US/10223711
 Publication No. US2003013344A1
 GENERAL INFORMATION:
 APPLICANT: Baralierz, Lauren O.
 APPLICANT: Kaumays, Pravin T.P.
 TITLE OF INVENTION: Synthetic Chimeric Filibrin Peptides

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; FILE REFERENCE: 18525/04058
; CURRENT APPLICATION NUMBER: US/10/223,711
; CURRENT FILING DATE: 2002-08-19
; PRIORITY NUMBER: 09/148,711
; PRIOR FILING DATE: 1998-09-04
; PRIORITY NUMBER: 08/460,502
; PRIOR FILING DATE: 1995-06-02
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 8
; LENGTH: 21
; TYPE: PRT
; ORGANISM: Clostridium tetani
US-10-223-711-8

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Query Match          60.2%;  Score 112;  DB 14;  Length 21;
Best Local Similarity 100.0%;  Pred. No. 3e-08;
Matches 21;  Conservative 0;  Mismatches 0;  Indels 0;  Gaps 0;
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Db      1 FNNFTVSPWLRYPKVSSASHLE 21

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RESULT 15
US-10-223-809A-6
; Sequence 6, Application US/10223809A
; Publication No. US20030157117A1
; GENERAL INFORMATION:
; APPLICANT: Pharmexa A/S
; ATTORNEY: Rasmussen, Peter Birk et al.
; TITLE OF INVENTION: No. US20030157117A1 Method for Down-Regulation of Amyloid
; FILE REFERENCE: 67542-2008
; CURRENT APPLICATION NUMBER: US/10/223,809A
; CURRENT FILING DATE: 2002-08-20
; PRIOR APPLICATION NUMBER: US 60/337,543
; PRIOR FILING DATE: 2001-08-20
; PRIOR APPLICATION NUMBER: US 60/373,027
; PRIOR FILING DATE: 2002-04-16
; PRIOR APPLICATION NUMBER: DE 2001 01231
; PRIOR FILING DATE: 2001-08-20
; PRIOR APPLICATION NUMBER: DE 2002 0058
; NUMBER OF SEQ ID NOS: 17
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 6
; LENGTH: 21
; TYPE: PRT
; ORGANISM: Clostridium tetani
US-10-223-809A-6

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Query Match          60.2%;  Score 112;  DB 14;  Length 21;
Best Local Similarity 100.0%;  Pred. No. 3e-08;
Matches 21;  Conservative 0;  Mismatches 0;  Indels 0;  Gaps 0;
Qy      1 FNNFTVSPWLRYPKVSSASHLE 21
Db      1 FNNFTVSPWLRYPKVSSASHLE 21

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Job time : 26.7237 secs

GenCore version 5.1.6
 Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: March 10, 2004, 08:58:54 ; Search time 33:6031 Seconds
 (without alignments)
 319.245 Million cell updates/sec

Title: US-09-848-834A-10
 Perfect score: 186

Scoring table: BLOSSOM62
 Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0

Maximum DB seq length: 20000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

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3: sp_fungi:*

4: sp_invertebrate:*

5: sp_mammal:*

6: sp_mhc:*

7: sp_mhc:*

8: sp_organelle:*

9: sp_phage:*

10: sp_plant:*

11: sp_rabbit:*

12: sp_virus:*

13: sp_vertebrate:*

14: sp_unclassified:*

15: sp_virus:*

16: sp_bacteriophage:*

17: sp_archeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	112	60.2	451	2 O9LA13	Osal13 clostridium
2	112	60.2	1310	2 Q93N7	Q93n27 clostridium
3	62	33.3	1268	2 Q45851	Q45851 clostridium
4	61	32.8	1278	2 Q57236	Q57236 clostridium
5	58	31.2	361	2 Q45848	Q45848 clostridium
6	58	31.2	361	2 Q45846	Q45846 clostridium
7	58	31.2	441	2 Q9X708	Q9x708 clostridium
8	58	31.2	1291	2 Q9ZAJ8	Q9zaj8 clostridium
9	58	31.2	1291	2 Q93G71	Q93g71 clostridium
10	58	31.2	1291	2 Q933K0	Q933k0 clostridium
11	58	31.2	1291	2 Q98077	Q98077 clostridium
12	58	31.2	1291	2 Q8GG96	Q8gr96 clostridium
13	57	30.6	430	2 Q9XAV1	Q9xav1 pseudomonas
14	57	30.6	502	16 Q98T8	Q9x8t8 streptomyces
15	56.5	30.4	367	2 Q45861	Q45861 clostridium
16	56.5	30.4	367	2 Q45862	Q45862 clostridium

Result No.	Score	Query Match	Length	DB ID	Description
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6	58	31.2	361	2 Q45846	Q45846 clostridium
7	58	31.2	441	2 Q9X708	Q9x708 clostridium
8	58	31.2	1291	2 Q9ZAJ8	Q9zaj8 clostridium
9	58	31.2	1291	2 Q93G71	Q93g71 clostridium
10	58	31.2	1291	2 Q933K0	Q933k0 clostridium
11	58	31.2	1291	2 Q98077	Q98077 clostridium
12	58	31.2	1291	2 Q8GG96	Q8gr96 clostridium
13	57	30.6	430	2 Q9XAV1	Q9xav1 pseudomonas
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6	58	31.2	361	2 Q45846	Q45846 clostridium
7	58	31.2	441	2 Q9X708	Q9x708 clostridium
8	58	31.2	1291	2 Q9ZAJ8	Q9zaj8 clostridium
9	58	31.2	1291	2 Q93G71	Q93g71 clostridium
10	58	31.2	1291	2 Q933K0	Q933k0 clostridium
11	58	31.2	1291	2 Q98077	Q98077 clostridium
12	58	31.2	1291	2 Q8GG96	Q8gr96 clostridium
13	57	30.6	430	2 Q9XAV1	Q9xav1 pseudomonas
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8	58	31.2	1291	2 Q9ZAJ8	Q9zaj8 clostridium
9	58	31.2	1291	2 Q93G71	Q93g71 clostridium
10	58	31.2	1291	2 Q933K0	Q933k0 clostridium
11	58	31.2	1291	2 Q98077	Q98077 clostridium
12	58	31.2	1291	2 Q8GG96	Q8gr96 clostridium
13	57	30.6	430	2 Q9XAV1	Q9xav1 pseudomonas
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16	56.5	30.4	367	2 Q45862	Q45862 clostridium

Result No.	Score	Query Match	Length	DB ID	Description
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6	58	31.2	361	2 Q45846	Q45846 clostridium
7	58	31.2	441	2 Q9X708	Q9x708 clostridium
8	58	31.2	1291	2 Q9ZAJ8	Q9zaj8 clostridium
9	58	31.2	1291	2 Q93G71	Q93g71 clostridium
10	58	31.2	1291	2 Q933K0	Q933k0 clostridium
11	58	31.2	1291	2 Q98077	Q98077 clostridium
12	58	31.2	1291	2 Q8GG96	Q8gr96 clostridium
13	57	30.6	430	2 Q9XAV1	Q9xav1 pseudomonas
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15	56.5	30.4	367	2 Q45861	Q45861 clostridium
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6	58	31.2	361	2 Q45846	Q45846 clostridium
7	58	31.2	441	2 Q9X708	Q9x708 clostridium
8	58	31.2	1291	2 Q9ZAJ8	Q9zaj8 clostridium
9	58	31.2	1291	2 Q93G71	Q93g71 clostridium
10	58	31.2	1291	2 Q933K0	Q933k0 clostridium
11	58	31.2	1291	2 Q98077	Q98077 clostridium
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5	58	31.2	361	2 Q45848	Q45848 clostridium
6	58	31.2	361	2 Q45846	Q45846 clostridium
7	58	31.2	441	2 Q9X708	Q9x708 clostridium
8	58	31.2	1291	2 Q9ZAJ8	Q9zaj8 clostridium
9	58	31.2	1291	2 Q93G71	Q93g71 clostridium
10	58	31.2	1291	2 Q933K0	Q933k0 clostridium
11	58	31.2	1291	2 Q98077	Q98077 clostridium
12	58	31.2	1291	2 Q8GG96	Q8gr96 clostridium
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3	62	33.3	1268	2 Q45851	Q45851 clostridium
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5	58	31.2	361	2 Q45848	Q45848 clostridium
6	58	31.2	361	2 Q45846	Q45846 clostridium
7	58	31.2	441	2 Q	

Q93N27	PRELIMINARY;	PRT;	1310 AA.	
Q93N27;				
AC				
01-DEC-2001	(TREMBLrel. 19, Created)			
01-DEC-2001	(TREMBLrel. 19, Last sequence update)			
DT				
01-OCT-2003	(TREMBLrel. 25, Last annotation update)			
DR				
Tetanus toxin (Fragment).				
Clostridium tetani.				
Bacteria; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;				
Clostridium				
NCBI_TaxID=1513;				
[1]	SEQUENCE FROM N.A.			
Shinmin Z., Dianling L.;	"Cloning and sequence analysis of tetanus toxin gene.";			
Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.				
EMBL; AFS89424; AAK72964.2;				
InterPro; IPR00237; Fendometallopeptidase inhibitor activity; IEA.				
GO; GO:0008237; Fendometallopeptidase activity; IEA.				
GO; GO:0015070; Zinc ion binding; IEA.				
GO; GO:008270; F-toxin activity; IEA.				
GO; GO:0009405; Pathogenesis; IEA.				
GO; GO:0006508; Peptidolysis and peptidolysis; IEA.				
InterPro; IPR001084; Cona1-like_ec_g1				
InterPro; IPR001064; Crystallin.				
InterPro; IPR002160; Kunzite, legume.				
InterPro; IPR000385; Peptidase_M27.				
InterPro; IPR006023; Peptidase_M2n_BS.				
Pfam; PF01742; Peptidase_M27_1.				
PRNTS; PR00760; BONTOXILYSIN.				
PRODOM; PD001963; Bontoxin type F.				
PROSITE; PS00225; CRYSTALLIN BETAGAMMA; 1.				
PROSITE; PS00142; ZINC_Protease; 1.				
NON-TER 1 1				
NON-TER 1310 1310				
SEQUENCE 1310 AA, 150316 MW; 9EADDCC914418E450 CRC64;				
Query Match Score 112; DB 2; Length 1310;				
Best Local Similarity 100.0%; Pred. No. 5e-07; Indels 0; Gaps 0;				
Matches 21; Conservative 0; Mismatches 0; RT				
Q94851	PRELIMINARY;	PRT;	1268 AA.	
AC				
01-NOV-1996 (TREMBLrel. 01, Created)				
01-NOV-1996 (TREMBLrel. 01, Last sequence update)				
DT				
01-OCT-2003 (TREMBLrel. 25, Last annotation update)				
DR				
Neurotoxin type F.				
BONT / F				
Clostridium baratii.				
Bacteria; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;				
Clostridium				
NCBI_TaxID=1561;				
[1]	SEQUENCE FROM N.A.			
Thompson D.E., Allaway D., Collins M.D.,				
Richardson P.T.;				
"Nucleotide sequence of the gene coding for Clostridium baratii type F neurotoxin: comparison with other clostridial neurotoxins.";				
EMBL; X68262; CAA48329.1; -.				
PIR; S33411; S33411.				
HSSP; P10845; 3BTA.				
MEROPS; M27.002; -.				
GO; GO:0004866; Endopeptidase inhibitor activity; IEA.				
GO; GO:0008237; Fendometallopeptidase activity; IEA.				
GO; GO:0015070; F-toxin activity; IEA.				
GO; GO:0009405; Zinc ion binding; IEA.				
GO; GO:0009406; Participation; IEA.				

DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE BonT Protein.
GN
OS Clostridium botulinum; Clostridia; Clostridiaceae;
OC Bacteria; Firmicutes; Clostridia; Clostridiaceae;
OC Clostridium.
OX NCBI_TaxID=1491;
RN [1] _
RP SEQUENCE FROM N.A.
RC STRAIN=CDC 3281;
RX MEDLINE=98440321; PubMed=3767710;
RA Santos-Buelga J., Collins M.D., East A.K.;
RT "Characterization of the genes encoding the Botulinum neurotoxin complex in a strain of clostridium botulinum producing type B & F neurotoxins.",
RL Curr. Microbiol. 37:312-318 (1998).
DR Y13630; CBA73968.1; -.
DR HSSP; P10845; 3PTA.
DR GO; GO:0008466; F:endopeptidase inhibitor activity; IEA.
DR GO; GO:0008377; F:metallopeptidase activity; IEA.
DR GO; GO:0015070; F:toxin activity; IEA.
DR GO; GO:00083270; F:zinc ion binding; IEA.
DR GO; GO:0009405; F:pathogenesis; IEA.
DR GO; GO:0006508; F:proteolysis and peptidolysis; IEA.
DR InterPro; IPR00985; ConA_like_lec_g1.
DR InterPro; IPR002160; Kunitz_1Legume.
DR InterPro; IPR00395; Pept_M_Zn_BS.
DR Pfam; PF01742; Peptidase_M27; 1.
DR PRINTS; PR00760; BONTOXILYSIN.
DR PRODOM; PD001963; Bontoxilysin. 1.
DR PROSITE; PS00142; ZINC_ProteaseB; 1.
SEQUENCE 1291 AA; 150840 MW; B4D3B0E46AB2E735 CRC64;

Query Match 31.2%; Score 58; DB 2; Length 1291;
Best Local Similarity 64.3%; Pred. No. 25;
Matches 9; Conservative 4; Mismatches 1; Indels 0; Gaps 0;
SQ

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1 FNNFTYSEFWLRVPK 14
Db 923 FLDFSVSEFWIRPK 936

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RESULT 9
ID Q93G71 PRELIMINARY; PRT; 1291 AA.
AC Q93G71;
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DR Clostridium botulinum.
OC Bacteria; Firmicutes; Clostridia; Clostridiaceae;
OC NCBI_TaxID=1491;
RN SEQUENCE FROM N.A.
RC STRAIN=1436;

RA Kirma N., Ferreira J.L., Baumstark B.R.;
RT "Characterization of six type A strains of Clostridium botulinum that contain type B toxin gene sequences";
RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF300466; AAL11499.1; -
DR EMBL; AF300455; AAL11498.1; -
DR GO; GO:0008466; F:endopeptidase inhibitor activity; IEA.
DR GO; GO:0008237; F:metallopeptidase activity; IEA.
DR GO; GO:0015070; F:toxin activity; IEA.
DR GO; GO:0008270; F:zinc ion binding; IEA.
DR GO; GO:0009405; F:pathogenesis; IEA.
DR InterPro; IPR00395; Peptidolysis and Peptidolysis; IEA.
DR InterPro; IPR003160; ConA_like_lec_g1.
DR InterPro; IPR00395; Peptidolysis; IEA.
DR InterPro; IPR00385; ConA_like_lec_g1.
DR InterPro; IPR003160; Kunitz_1Legume.
DR InterPro; IPF01742; Peptidase_M27; 1.
DR PRINTS; PR00760; BONTOXILYSIN.
DR PRODOM; PD001963; Bontoxilysin. 1.
DR PROSITE; PS00142; ZINC_ProteaseB.
SQ SEQUENCE 1291 AA; 150843 MW; TAC1737B0PASA151 CRC64;

Query Match 31.2%; Score 58; DB 2; Length 1291;
Best Local Similarity 64.3%; Pred. No. 25;
Matches 9; Conservative 4; Mismatches 1; Indels 0; Gaps 0;
SQ

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1 FNNFTYSEFWLRVPK 14
Db 923 FLDFSVSEFWIRPK 936

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RESULT 11
ID Q08077 PRELIMINARY; PRT; 1291 AA.
AC Q08077;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DR BONT/B.
GN Clostridium botulinum.
OC Bacteria; Firmicutes; Clostridia; Clostridiaceae;

OC Clostridium
NCBI_TaxID=1491;
OX [1]
RN SEQUENCE FROM N.A.
STRAIN=Ex1lund 17B ATCC25765;
RC DR GO:0008237; F:metalloendopeptidase inhibitor activity; IEA.
RX DR GO:0015070; F:toxin activity; IEA.
RA Hutson R.A., Collins M.D., East A.K., Thompson D.E.;
RT "Nucleotide sequence of the gene coding for non-proteolytic
clostridial neurotoxins."
RT Clostridium botulinum type B neurotoxin: comparison with other
Cllr. Microbiol. 28:101-110(1994).
RL EMBL; X71343; CAJ50482.1;
DR PIR; I0631; 140631.
HSSP; P10845; 3BTA.
DR MEROPS; M27.002;
DR GO; GO:0004866; F:endopeptidase inhibitor activity; IEA.
DR GO; GO:0015070; F:toxin activity; IEA.
DR GO; GO:0008270; F:inc ion binding; IEA.
DR GO; GO:0009405; P:Pathogenesis; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR008905; ConA-like_lec_g1.
DR InterPro; IPR002160; Kunitz_legume.
DR InterPro; IPR00035; Peptidase_M27.
DR InterPro; IPR006025; Pept_M_Zn_BS.
DR Pfam; PF001742; Peptidase_M27_1.
DR PRINTS; PR00760; BONTOXILYSIN.
DR ProDom; PD001963; Bontoxilysin; 1.
DR PROSITE; PS00142; ZINC PROTEASE; 1.
RN SEQUENCE 1291 AA; 150513 MW; 71B0CAFE23D69FAAA CRC64;
SQ [1]
Query Match Score 58 / DB 2; Length 1291;
Best Local Similarity 64.3%; Pred. No. 25;
Matches 9; Conservative 4; Mismatches 1; Indels 0; Gaps 0;
QY 1 FNNFTVSPMLRVPK 14
Db 923 FLDPVSFVWIRPK 936
RESULT 12
Q8GR96 ID Q8GR96 PRELIMINARY; PRT; 1291 AA.
AC [1]
RN SEQUENCE FROM N.A.
RA Ihara H., Kohda T., Morimoto F., Tsukamoto K., Karabawa T.,
Nakamura S., Mukamoto M., Kozaiki S.; Clostridium botulinum type B neurotoxin associated with infant
botulism";
RT Clostridium botulinum type B neurotoxin associated with infant
botulism";
EMBL; AB084152; BAC22064.1;
DR GO; GO:0004866; F:metalloendopeptidase inhibitor activity; IEA.
DR GO; GO:000237; F:metallopeptidase activity; IEA.
DR GO; GO:0015070; F:toxin activity; IEA.
DR GO; GO:0008270; F:inc ion binding; IEA.
DR GO; GO:0009405; P:Pathogenesis; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR008985; ConA-like_lec_g1.
DR InterPro; IPR002160; Kunitz_legume.
DR Pfam; PF001742; Pept_M_Zn_BS.
DR PRINTS; PR00760; BONTOXILYSIN.
DR PRODOM; PD001963; Bontoxilysin; 1.
DR PROSITE; PS00142; ZINC PROTEASE; 1.
SQ SEQUENCE 1291 AA; 150513 MW; 71B0CAFE23D69FAAA CRC64;
Query Match Score 58 / DB 2; Length 1291;
Best Local Similarity 64.3%; Pred. No. 25;
Matches 9; Conservative 4; Mismatches 1; Indels 0; Gaps 0;
QY 1 FNNFTVSPMLRVPK 14
Db 923 FLDPVSFVWIRPK 936
RESULT 13
Q9XAV1 ID Q9XAV1 PRELIMINARY; PRT; 430 AA.
AC [1]
RN SEQUENCE FROM N.A.
RA Q9XAV1 PRELIMINARY; PRT; 430 AA.
DR 01-NOV-1999 (TREMBLrel. 12, Created)
DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)
DE Alkane 1-monooxygenase (EC 1.14.15.1).
GN ALKB
PSA
Pseudomonas fluorescens.
Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
Pseudomonadaceae; Pseudomonas.
OC
OC
OC
NCBI_TaxID=294;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CHA0;
RA Submitted (MAR-2001) to the EMBL/GenBank/DDBJ databases.
DR STRAIN=CHA0;
RX MEDLINE=11207749; PubMed=11207749;
RA Smits T.H.M., Roethlisberger M., Van Beilen J.B.,
RT "Molecular screening for alkane hydroxylase genes in Gram-negative and
Gram-positive bacteria";
RL Environ. Microbiol. 1:307-317(1999).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=CHA0;
RA Smits T.H.M.;
DR EMBL; AT009539; CRB5105.2;
DR GO; GO:0018638; Ficamphor 5-monooxygenase activity; IEA.
DR InterPro; IPR005804; Ficamphor 5-monooxygenase activity; IEA.
DR Pfam; PF00487; FA_desat_fam.
DR Monooxygenase; Oxdoreductase;
KW SEQUENCE 430 AA; 48337 MW; 048950980783E86 CRC64;
SQ [1]
Query Match Score 57 / DB 2; Length 430;
Best Local Similarity 35.9%; Pred. No. 10;
Matches 14; Conservative 4; Mismatches 7; Indels 14; Gaps 2;
QY 7 SFWSFLPRTWFSLSAWHESQERLEKLGLPTLHKWNGY 262
Db 224 SFWSFLPRTWFSLSAWHESQERLEKLGLPTLHKWNGY 262
RESULT 14
Q9XBT8 ID Q9XBT8 PRELIMINARY; PRT; 502 AA.
AC [1]
RN SEQUENCE FROM N.A.
RA Ihara H., Kohda T., Morimoto F., Tsukamoto K., Karabawa T.,
Nakamura S., Mukamoto M., Kozaiki S.; Clostridium botulinum type B neurotoxin associated with infant
botulism";
RT Clostridium botulinum type B neurotoxin associated with infant
botulism";
EMBL; AB084152; BAC22064.1;
DR GO; GO:0004866; F:metalloendopeptidase inhibitor activity; IEA.
DR GO; GO:000237; F:metallopeptidase activity; IEA.
DR GO; GO:0015070; F:toxin activity; IEA.
DR GO; GO:0008270; F:inc ion binding; IEA.
DR GO; GO:0009405; P:Pathogenesis; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR008985; ConA-like_lec_g1.
DR InterPro; IPR002160; Kunitz_legume.
DR Pfam; PF001742; Pept_M_Zn_BS.
DR PRINTS; PR00760; BONTOXILYSIN.
DR PRODOM; PD001963; Bontoxilysin; 1.
DR PROSITE; PS00142; ZINC PROTEASE; 1.
SQ SEQUENCE FROM N.A.
RC STRAIN=A3 (2) / MI45;
RX MEDLINE=21996410; PubMed=12000953;
RA Bentley S.D., Chater K.F., Cerdeno-Tarraga A.-M., Challis G.L.,

RA Thomson N.R., James K.D., Harris D.E., Quail M.A., Kieser H.,
 RA Harper D., Bateman A., Brown S., Chaudra G., Chen C.W., Collins M.,
 RA Cronin A., Horrabin A., Hobday J., Hidalgo J., Howard S.,
 RA Huang C.-H., Kieser H., Goble A., Murphy L., Oliver K., O'Neil S.,
 RA Rabinowitsch E., Rajandream M.A., Rutherford K., Rutter S.,
 RA Seeger K., Saunders D., Sharp S., Squares R., Taylor K.,
 RA Warren T., Wierzorek A., Woodward J., Barrell B.G., Parthill J.,
 RA Hopwood D.A.;
 RT "Complete Genome sequence of the model actinomycete Streptomyces
 coelicolor A3 (2)" ;
 RT Nature 417:141-147 (2002);
 RL EMBL; AL931118; CAB32730.1; -.
 DR PIR; T36589; T36589.
 DR GO; GO:001021; C: integral membrane; IEA.
 KW Transmembrane; Complete proteome.
 SQ SEQUENCE 502 AA; 54795 MW; C84F774COA5AADBC CRC64;
 SEQUENCE 502 AA; 54795 MW; C84F774COA5AADBC CRC64;

Query Match 30.6%; Score 57; DB 16; Length 502;
 Best Local Similarity 50.0%; Pred. No. 12;
 Matches 8; Conservative 3; Mismatches 5; Indels 0; Gaps 0;
 QY 18 SHLEGPLHEVSYGLRP 33
 DB 483 AHFEGPQVHWGKGRQP 498

RESULT 15

Q45861 PRELIMINARY; PRT; 367 AA.
 ID Q45861.
 AC Q45861.
 DT 01-NOV-1996 (T-EMBLrel. 01, Created)
 DT 01-NOV-1996 (T-EMBLrel. 01, Last sequence update)
 DT 01-OCT-2003 (T-EMBLrel. 25, Last annotation update)
 DE Botulinum neurotoxin type B (Fragment).
 BONT_E.
 GN Clostridium botulinum.
 OS Clostridium botulinum.
 OC Bacteria; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;
 OC Clostridium.
 CX NCBI_TaxID:1491;
 PN [1]
 RP SEQUENCE FROM N. A.
 RC STRAIN=type E;
 RX MEDLINE=94013372; PubMed=8408542;
 RA Campbell K., East A.K., Collins M.D.;
 RT Gene probes for identification of the botulinal neurotoxin gene and
 specific identification of neurotoxin types B, E, and F. n;
 RL J. Clin. Microbiol. 31:2255-2262 (1993).
 DR EMBL; X70816; CA50149.1/-.
 DR PIR; S48106; S41106.
 DR HSSP; P10845; 3BTA.
 DR InterPro; IPR00985; ConA_like_lec_91.
 DR GO; GO:0015070; P:toxin activity; IEA.
 KW Neurotoxin.
 FT NON_TER 1 367 AA; 42902 MW; 346A610C2FF70262 CRC64;
 FT NON_TER 1 367 AA; 42902 MW; 346A610C2FF70262 CRC64;

Query Match 30.4%; Score 56.5; DB 2; Length 367;
 Best Local Similarity 22.4%; Pred. No. 9.9;
 Matches 15; Conservative 8; Mismatches 7; Indels 37; Gaps 2;
 SP 23
 QY 1 FNNEFTVSEFWLRVP-----KVSASHLE-----
 DB 297 YKNPSISFWRIPNYDNKIVNNEYIINCMDNNSGWYSLNHNELIMLQDNAGINQ 356

QY 24 SIEHWSYG 30
 DB 357 KLAFLNYIG 363

Search completed: March 10, 2004, 09:25:33
 Job time : 35.6031 secs

PA	(WANG/) WANG C. Y.	XX	PR 05-OCT-1998;	98DX-000001261.
PA	(ZAMB/) ZAMB T.	XX	PR 20-OCT-1998;	98US-01501P.
PI	Ladd AE, Wang CY, Zamb T;	XX	XX	
DR	WPI; 1994-357910/44.	XX	PA (MEBI-) M & E BIOTECH AS.	
XX	Immunogenic luteinising hormone releasing hormone peptide(s) - that suppress LHRH activity in males and females.	XX	PI Steinaa L, Mouritsen S, Nielsen KG, Haaning J, Leach D, Dalum I;	
PT	PT	XX	PI Gaturam A, Birk P, Karlsson G;	
XX	XX	XX	XX	WPI; 2000-345917/30.
PS	Claim 8; Page 84; 213pp; English.	XX	XX	DR
XX	Synthetic immunogenic peptides are provided in which a universal immune stimulator is linked to a peptide or protein hapten containing B cell and/or cytotoxic T lymphocyte epitope, giving a product which causes potent immune responses to the coupled peptide or protein. The stimulator consists of (A) a promiscuous helper T cell epitope (Th) which elicits an immune response to the coupled peptide in members of a heterogeneous population expressing diverse HLA phenotypes, and (B) an adjuvant peptide sequence from the invasin protein of Yersinia. Spacer amino acid sequences (e.g. Gly-Gly) can be provided between the invasin and Th domain and between the immune stimulator and hapten components. When the hapten is LHRH, then optionally the invasin domain can be omitted from the immune stimulator component. The present sequence represents an LHRH-containing, invasin-free immunogenic peptide as above which can be used as a potent vaccine for treating e.g. prostatic hyperplasia, androgen-dependent carcinoma, prostatic carcinoma, testicular carcinoma, endometriosis, benign uterine tumours, or oestrogen-dependent breast cancer, (severe) premenstrual syndrome or oestrogen-dependent breast cancer, or for induction of infertility. (Updated on 25-MAR-2003 to correct PN field.)	CC	XX	XX
XX	Sequence 32 AA;	CC	XX	XX
Query Match	Score 81.7%; Best Local Similarity 87.9%; Matches 29;	DB 2;	Score 152%; Pred. No. 5; 6e-15;	Length 32;
Best Local Similarity	Conservative 0;	Indels 0;	Mismatches 0;	
Matches				
Oy	1 FNNFTVFSMLRVPKVASHLLEGPSLHNSYGLRP 33	Db	1 FNNFTVFSMLRVPKVASHLLEGPSLHNSYGLRP 33	Db 673 FNNFTVFSMLRVPKVASHLLEGPSLHNSYGLRP 33
Db	3 FNNFTVFSMLRVPKVASHL---HNSYGLRP 31	XX	XX	Sequence 750 AA;
RESULT 4	AAV92633 standard; protein: 750 AA.	XX	XX	Query Match 65.3%; Score 121.5%; Length 750;
ID	AAV92633	AC	AC	Best Local Similarity 86.2%; Pred. No. 6.5e-09;
XX	AAV92633;	XX	XX	Matches 25; Conservative 0; Mismatches 3;
DT	10-AUG-2000 (first entry)	XX	XX	Indels 1; Gaps 1;
XX	Mutant human prostate specific membrane antigen construct, hPSM1.10.	XX	XX	Result 5
XX	Prostate specific membrane antigen; immunogenized construct; mutant; vaccination; cytotoxic T-lymphocyte immunity; breast cancer;	DE ABR82481	DE ABR82481 standard; protein: 537 AA.	
KW	prostate cancer; cell-associated peptide antigen; foreign epitope.	XX	XX	ID ABR82481
XX	Homo sapiens.	AC ABR82481;	AC ABR82481;	
OS	Synthetic.	XX	XX	
XX	Location/Qualifiers	XX	XX	Truncated human CEA-TT P2 and P30 epitopes.
FH	24..38	FH	XX	CEA; immune response; carinoembryonic antigen; antigen presenting cell;
Key Peptide	/label= P2	FT	AC	APC; cytostatic; vaccine; human; tetanus toxoid; p2; p30; antigen.
FT	/note= "foreign epitope"	FT	XX	XX
FT	673..693	FT	OS Synthetic.	
Peptide	/label= P30	FT	XX	Location/Qualifiers
FT	/note= "foreign epitope"	FT	XX	1..34
XX	WO200020027-A2.	FT	XX	/note= "signal peptide"
XX	XX	FT	35..537	35..537
PN	PN	FT	XX	/note= "mature protein"
PD	13-APR-2000.	XX	XX	WO2003059379-A2.
XX	05-OCT-1999;	XX	XX	XX

PD 24-JUL-2003.
 XX
 PF 17-JAN-2003; 2003WO-DK000031.
 XX
 PR 17-JAN-2002; 2002DK-0000082.
 PR 17-JAN-2002; 2002US-0350047P.
 XX
 PA (PHAR-) PHARMEXA AS.
 XX
 PI Klysnar S, Voldborg B;
 XX
 WPI; 2003-587260/55.
 DR N-PSDB; ACF35966.
 XX
 PT Inducing an immune response in humans against autologous carcinoembryonic antigen (CEA) comprising administering a modified CEA polypeptide, a nucleic acid encoding the polypeptide, or a microorganism expressing the polypeptide.
 XX
 Disclosure; Page 121-124; 140PP; English.
 PS
 PT Inducing an immune response against autologous carcinoembryonic antigen (CEA) in an animal, e.g. human. The method involves effecting uptake and processing by antigen presenting cells (APCs) in the animal of at least 1 modified CEA polypeptide or of a nucleic acid encoding the modified CEA polypeptide or of a microorganism or virus expressing the modified CEA polypeptide to induce a CTL response and an antibody response that targets the autologous CEA. The method is useful in immunizing actively against diseases characterized by cells that express CEA. The present sequence represents a modified human CEA polypeptide that has tetanus toxoid (TT) P2 and P30 epitopes introduced in its sequence
 XX
 Sequence 537 AA;
 SQ
 Query Match 65.1%; Score 121; DB 7; Length 708;
 Best Local Similarity 95.8%; Pred. No. 5.3e-09;
 Matches 23; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 XX
 Sequence 537 AA;
 SQ
 Query Match 65.1%; Score 121; DB 7; Length 537;
 Best Local Similarity 95.8%; Pred. No. 5.3e-09;
 Matches 23; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 XX
 Query 1 FNNFTVFWLRPKVVSASHLTPQHQ 667
 DB 513 ENNFTVFWLRPKVVSASHLTPQHQ 536
 RESULT 7
 ABR82478
 ID ABR82478 standard; protein; 717 AA.
 XX
 AC ABR82478;
 XX
 DT 20-NOV-2003 (first entry)
 XX
 DE Modified human CEA-TT P2 and P30 epitopes.
 XX
 CA ABR82479;
 XX
 DT 20-NOV-2003 (first entry)
 XX
 DE Modified human CEA-TT P2 and P30 epitopes.
 XX
 KW CBA; immune response; carcinoembryonic antigen; antigen presenting cell;
 KW APC; cytostatic; vaccine; human; tetanus toxoid; p2; p30; antigen.
 XX
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 Peptide 1 .34
 FT /note= "signal peptide"
 FT 35 .708
 Protein /note= "mature protein"
 FT
 XX WO2003059379-A2.
 PN
 XX
 PD 24-JUL-2003.
 XX
 FT 17-JAN-2002; 2002DK-0000082.
 PR 17-JAN-2002; 2002US-0350047P.
 XX
 PA (PHAR-) PHARMEXA AS.
 XX
 PI Klysnar S, Voldborg B,
 XX
 WPI; 2003-587260/55.

DR N-PSDB; ACF35964.
 XX Inducing an immune response in humans against autologous carcinoommibronic
 PT antigen (CEA) comprises administering a modified CEA polypeptide, a
 PT nucleic acid encoding the polypeptide, or a microorganism expressing the
 PT polypeptide.
 XX Disclosure; Page 114-117; 140pp; English.
 PS The invention relates to inducing an immune response against autologous
 CC carcinembryonic antigen (CEA) in an animal, e.g. human. The method
 CC involves effecting uptake and processing by antigen presenting cells
 CC (APCs) in the animal of at least 1 modified CEA polypeptide or a
 CC nucleic acid encoding the modified CEA polypeptide or of a microorganism
 CC or virus expressing the modified CEA polypeptide to induce a CTL response
 CC and an antibody response that targets the autologous CEA. The method is
 CC useful in immunizing actively against diseases characterized by cells
 CC that express CEA. The present sequence represents a modified human CEA
 CC polypeptide that has tetanus toxoid (TT) P2 and P30 epitopes introduced
 CC in its sequence
 XX Sequence 717 AA;
 SQ Query Match 65.1%; Score 121; DB 717; Length 717;
 Best Local Similarity 95.8%; Pred. No. 7 4e-09; 1; Indels 0; Gaps 0;
 Matches 23; Conservative 0; Mismatches 1; Delins 0; Gaps 0;
 Qy 1 FNNFTVSFMLRVPKVVASLLEGPS 24
 Db 693 FNNFTVSFMLRVPKVVASLLEGTS 716

RESULT 8
AAO30488 standard; protein; 194 AA.

ID AAO30488
 XX AC
 XX DT 22-SEP-2003 (first entry)
 DE Human TNFalpha variant, TNF24-P2-P30.

XX Multimeric protein, interleukin 5; IL5; TNFalpha; inflammatory disease;
 KW tumour necrosis factor alpha; gene therapy; arthritis; human; mutant;
 KW mutant; variant; tetanus toxoid; epitope.
 XX Homo sapiens.
 OS Unidentified.
 OS Chimeric.
 XX Key Location/Qualifiers
 FH 2-.109
 FT /note= "Human TNF"
 FT Region
 FT 110-.124
 FT /note= "Tetanus toxoid P2 epitope"
 FT Region
 FT 125-.145
 FT /note= "Tetanus toxoid P30 epitope"
 FT Region
 FT 146-.194
 FT /note= "Human TNF"
 XX WO200302244-A2.
 XX PD 22-MAY-2003.
 XX PF 15-NOV-2002; 2002NO-DK000764.
 XX PR 16-NOV-2001; 2001DX-00001702.
 PR 16-NOV-2001; 2001US-0331575P.
 XX PA (PHAR-) PHARMAXA AS.
 PA (KLYS/) KLYSNER S.
 PA (NIEL/) NIELSEN F. S.
 PA (BRAT/) BRATT T.

PA (VOLD/) VOLDBORG B.
 PA (MOUR/) MOURITSEN S.
 PA XX Klysnar S, Nielsen FS, Bratt T, Voldborg B, Mouritzen S,
 PI XX
 XX DR WPI; 2003-44558/42.
 XX PT New immunogenic analogue of a polymeric protein, useful for preparing a
 PT composition for treating inflammatory diseases e.g. arthritis.
 XX
 PS Claim 23; Page 158; 196pp; English.
 XX
 CC The invention relates to immunogenic analogues of multimeric proteins
 CC such as immunogenic variants of interleukin 5 (IL5) and tumour necrosis
 CC factor alpha (TNF, TNFalpha), and methods for production of immunogenic
 CC analogues. The immunogenic analogue is useful for preparing a composition
 CC for treating inflammatory diseases, e.g., arthritis. It is also used in
 CC gene therapy. The present sequence is human TNFalpha variant protein with
 CC an inserted tetanus toxoid P2 and P30 epitopes. This sequence is used to
 CC illustrate the method of the invention
 XX
 SQ Sequence 194 AA;
 SQ Query Match 64.0%; Score 119; DB 6; Length 194;
 Best Local Similarity 81.5%; Pred. No. 3 2e-09; 4; Mismatches 1; Indels 0; Gaps 0;
 Matches 22; Conservative 4; Delins 0; Gaps 0;
 Qy 1 FNNFTVSFMLRVPKVVASLLEGPSHW 27
 Db 125 FNNFTVSFMLRVPKVVASLLEAZAKPW 151

RESULT 9
AAY92655 standard; peptide; 31 AA.

ID AAY92655
 XX AC
 XX DT 10-AUG-2000 (first entry)
 DE DE PSMpep012 - P30 inserted in hPSM insertion position 10.
 XX
 KW Foreign epitope; P2; prostate specific membrane antigen; vaccination;
 KW cytotoxic T-lymphocyte immunity; self-protein; cancer; breast cancer;
 KW prostate cancer; cell-associated peptide antigen.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX PH Key Location/Qualifiers
 FT 6-.26
 FT Peptide /label= P30
 XX PR WO200020027-A2.
 XX PD 13-APR-2000.
 XX PF 05-OCT-1999; 99WO-DK000525.
 XX PR 05-OCT-1998; 98DK-00001261.
 XX PR 20-OCT-1998; 98US-0105011P.
 XX PA (MEBI-) M & E BICTECH AS.
 XX PR 05-OCT-1998; 98DK-00001261.
 XX PA 20-OCT-1998; 98US-0105011P.
 XX PA (MEBI-) M & E BICTECH AS.
 XX PI Steinna L, Mouritzen S, Nielsen KG, Haanning J, Leach D, Dalum I;
 PI Gautam A, Birk P, Karlsson G,
 XX DR WPI; 2000-349917/30.
 XX PT Inducing immune responses to weakly immunogenic, tumor associated peptide
 PT antigens for the treatment of breast and prostate cancer.
 XX Example 1; Page 118; 220pp; English.
 PS

XX AAY92650-55 are peptides designed which correspond to the P2 and P3⁰ CC specific membrane antigen (hPSM) CC epitopes with 5 flanking amino acids correspond to the CC amino acids in each end. The flanking amino acids will be used in, e.g. T CC epitope insertion sites 6, 8 and 10. The peptides will be used in vitro assays, CC but also for ELISA or other immunological methods for inducing immune responses against weakly CC immunogenic cell-associated peptide antigens (e.g. those associated with cancers (i.e. self-proteins), for example, hPSM, CC heregulin 2 (Her2) and/or fibroblast growth factor 8b (FGF8b). The method CC comprises effecting simultaneous presentation by antigen producing cells (APCs) of the animals immune system of: (1) at least 1 CTL (cytotoxic T-CC lymphocyte) group derived from the PA and/or at least 1 B-cell helper CC derived from the cell-associated PA; and (2) at least 1 first T helper CC cell group which is foreign to the animal. Analogues of human PSM, human CC Her2 and human/murine FGF8b comprising a substantial part of all known CC and predicted CTL and B-cell epitopes of the respective PA and including CC at least one foreign T helper epitope (e.g. P2 and/or P3⁰) are also CC claimed. The method is used to treat prostate, prostate/breast or breast CC cancer when the PA is human PSM, FGF8b and Her2, respectively XX

Sequence 31 AA;

Query	1 FNNFTYSEWLVPKVSASHLEGPSLH	Score 118.5;	DB 2;	Length 31;
Best Local Similarity	92.3%	Pred. Nc.	4.6e-10;	
Matches	24;	Mismatches	0;	
Db	6 FNNFTYSEWLVPKVSASHLE-PSSH	1;	Indels	1;
			Gaps	1;

RESULT 10

AAW81336 standard; protein; 158 AA.

XX	AAW81336;	AC	AAW81336;	DT	21-APR-1999 (first entry)
XX				DE	TNF30-5, a TNF-alpha analogue.
XX				XX	Human tumour necrosis factor-alpha; TNF-alpha; TNF-alpha analogue; vaccine; rheumatoid arthritis; Crohn's disease; ulcerative colitis; cancer; disseminated sclerosis; diabetes; psoriasis; osteoporosis; asthma.
XX				OS	Synthetic.
XX				OS	Homo sapiens.
XX				PN	WO9846642-A1.
XX				PD	22-OCT-1998.
XX				PP	15-APR-1998; 98WO-DK000157.
XX				PR	15-APR-1997; 97DK-00000418.
XX				PR	24-APR-1997; 97US-0044187P.
XX				PA	(FERR) FARM LAB FERRING AS.
XX				PI	Jensen MR, Mouritsen S, Elsner H, Dalmat;
XX				DR	WPI, 1998 594561/50.
XX				DR	N-PSDB; AA968425.

XX				XX	Modified human tumour necrosis factor-alpha - comprises immunodominant T cell epitope, useful in vaccines to treat or prevent TNF-associated diseases, e.g. cancer.
XX				XX	Claim 15; Page 81-92; 134DP; English.
XX				XX	The present sequence represents a modified human tumour necrosis factor-

alpha (TNF-alpha) analogue. The analogues have no residual TNF activity CC and are immunogenic in a large number of the human population (by CC using promiscuous epitopes). The TNF-alpha analogue is able to generate, CC in humans, neutralizing antibodies to wild-type human TNF alpha has at CC least one fragment of TNF substituted by a peptide containing an CC immunodominant T-cell epitope, and at least one TNF-alpha B-cell epitope. CC The substitution causes a significant change in the amino acid sequence CC of any one of the strands in the front beta-sheet, any of the connecting CC loops or any of the B', I or D strands in the back beta-sheet. The TNF- CC alpha analogues are used as vaccines for treatment or prevention of CC diseases associated with excessive release or activity of TNF-alpha, e.g. CC rheumatoid arthritis, Crohn's disease, ulcerative colitis, cancer of any CC sort, disseminated sclerosis, diabetes, psoriasis, osteoporosis and CC asthma XX

Sequence 158 AA;

Query	1 FNNFTYSEWLVPKVSASHLEG 22	Score 118;	DB 2;	Length 158;
Best Local Similarity	100.0%;	Pred. No.	3.0e-09;	
Matches	22;	Conservative	0;	Mismatches 0;
Db	133 FNNFTYSEWLVPKVSASHLEG 154	Indels	0;	Gaps 0;

RESULT 11

Query	1 FNNFTYSEWLVPKVSASHLEG 22	Score 118.5;	DB 2;	Length 158 AA.
Best Local Similarity	92.3%;	Pred. Nc.	4.6e-10;	
Matches	24;	Mismatches	0;	
Db	133 FNNFTYSEWLVPKVSASHLEG 154	Indels	0;	

RESULT 12

Query	1 FNNFTYSEWLVPKVSASHLEG 22	Score 118.5;	DB 2;	Length 158 AA.
Best Local Similarity	92.3%;	Pred. Nc.	4.6e-10;	
Matches	24;	Mismatches	0;	
Db	133 FNNFTYSEWLVPKVSASHLEG 154	Indels	0;	

RESULT 13

Query	1 FNNFTYSEWLVPKVSASHLEG 22	Score 118.5;	DB 2;	Length 158 AA.
Best Local Similarity	92.3%;	Pred. Nc.	4.6e-10;	
Matches	24;	Mismatches	0;	
Db	133 FNNFTYSEWLVPKVSASHLEG 154	Indels	0;	

RESULT 14

Query	1 FNNFTYSEWLVPKVSASHLEG 22	Score 118.5;	DB 2;	Length 158 AA.
Best Local Similarity	92.3%;	Pred. Nc.	4.6e-10;	
Matches	24;	Mismatches	0;	
Db	133 FNNFTYSEWLVPKVSASHLEG 154	Indels	0;	

RESULT 15

Query	1 FNNFTYSEWLVPKVSASHLEG 22	Score 118.5;	DB 2;	Length 158 AA.
Best Local Similarity	92.3%;	Pred. Nc.	4.6e-10;	
Matches	24;	Mismatches	0;	
Db	133 FNNFTYSEWLVPKVSASHLEG 154	Indels	0;	

RESULT 16

Query	1 FNNFTYSEWLVPKVSASHLEG 22	Score 118.5;	DB 2;	Length 158 AA.
Best Local Similarity	92.3%;	Pred. Nc.	4.6e-10;	
Matches	24;	Mismatches	0;	
Db	133 FNNFTYSEWLVPKVSASHLEG 154	Indels	0;	

RESULT 17

Query	1 FNNFTYSEWLVPKVSASHLEG 22	Score 118.5;	DB 2;	Length 158 AA.
Best Local Similarity	92.3%;	Pred. Nc.	4.6e-10;	
Matches	24;	Mismatches	0;	
Db	133 FNNFTYSEWLVPKVSASHLEG 154	Indels	0;	

The present sequence represents a modified human tumour necrosis factor-

XX				XX	The invention provides a pharmaceutical vaccine composition (I) for the CC prevention or treatment of a self-protein-mediated pathology. The CC composition comprises a last one modified immunogenic self-protein CC (selected from modified TNF-alpha proteins) and a surfactant capable of CC acting as a solubilizer. (I) is useful for preventing or treating a self CC -protein-mediated pathology such as an inflammatory disease, rheumatoid CC arthritis, an inflammatory bowel disease (ulcerative colitis or Crohn's
----	--	--	--	----	---

aspirin).

Claim 21; Page 42-43; 55DP; English.

XX				XX	The invention provides a pharmaceutical vaccine composition (I) for the CC prevention or treatment of a self-protein-mediated pathology. The CC composition comprises a last one modified immunogenic self-protein CC (selected from modified TNF-alpha proteins) and a surfactant capable of CC acting as a solubilizer. (I) is useful for preventing or treating a self CC -protein-mediated pathology such as an inflammatory disease, rheumatoid CC arthritis, an inflammatory bowel disease (ulcerative colitis or Crohn's
----	--	--	--	----	---

aspirin).

Claim 22; Page 42-43; 55DP; English.

XX				XX	The invention provides a pharmaceutical vaccine composition (I) for the CC prevention or treatment of a self-protein-mediated pathology. The CC composition comprises a last one modified immunogenic self-protein CC (selected from modified TNF-alpha proteins) and a surfactant capable of CC acting as a solubilizer. (I) is useful for preventing or treating a self CC -protein-mediated pathology such as an inflammatory disease, rheumatoid CC arthritis, an inflammatory bowel disease (ulcerative colitis or Crohn's
----	--	--	--	----	---

aspirin).

CC disease), cancer, cachexia, multiple sclerosis, diabetes, psoriasis, CC osteoporosis or asthma. (1) is useful for inducing autoantibodies to a CC self-protein such as TNF (tumour necrosis factor)-alpha in a human CC subject. (1) comprising cetylpyridinium chloride as a component is useful CC for immunisation of a human subject or for treatment of a human CC inflammatory disease. The present sequence represents a human TNF-alpha CC analogue TNF30-5

XX Sequence 158 AA;

Query Match 63.4%; Score 118; DB 5; Length 158;
Best Local Similarity 100.0%; Pred. No. 3.6e-09;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 FNNFTVSEWLRLPKVSSASHLLEG 22

Db 133 FNNFTVSEWLRLPKVSSASHLLEG 154

RESULT 12

ID AAY92627 standard; protein; 750 AA.

XX AAY92627;

XX DT 10-AUG-2000 (first entry)

XX Mutant human prostate specific membrane antigen construct, hPSM1.1.

DE Prostate specific membrane antigen; immunogenized construct; mutant;
KW vaccination; cytotoxic T-lymphocyte immunity; breast cancer;
KW prostate cancer; cell-associated peptide antigen; foreign epitope.
XX

OS Homo sapiens.

XX Synthetic.

FH Location/Qualifiers

FT Peptide 17..31

/note= "foreign epitope"
FT /label= P2

FT Peptide 32..52

/note= "foreign epitope"
FT /label= P30

FT /note= "foreign epitope"

XX WO2000020027-A2.

PN PR 20-OCT-1998; 9BDK-00001261.

XX PR 20-OCT-1998; 9BSN-0105011P.

XX PA (MBI-) M & E BIOTECH AS.

XX PI Steinnaa L, Mouritsen S, Nielsen KG,

PI Gautam A, Birk P, Karlsson G;

XX DR XX

XX WPI; 2000-349917/30.

XX PT Inducing immune responses to weakly immunogenic, tumor associated peptide
PR antigens for the treatment of breast and prostate cancer.
XX Example 1; Page; 220pp; English.

PS XX

CC AAY92627-49 are mutant immunogenized human prostate specific membrane CC antigen (PSM) constructs, which contain foreign epitopes (P2 and/or P30). CC The immunogenic analogues of PSM can be used in the claimed method as an CC adjuvaccine to induce a CTL response. Subdominant CTL epitopes, antibody CC binding regions and cysteine residues involved in disulfide bonds are CC preserved in the immunogenized forms. The method is used for inducing CC immune responses against weakly immunogenic cell-associated peptide CC antigens (PA) such as those associated with cancers (self-proteins), e.g.

CC human prostate specific membrane antigen (PSM), heregulin 2 (Her2) and/or CC fibroblast growth factor 8b (FGF8b). The method comprises effecting CC simultaneous presentation by antigen producing cells (APCs) of the CC animals immune system of: (1) at least 1 CTL (cytotoxic T-lymphocyte) CC group derived from the PA and/or at least 1 B-cell group derived from the CC cell-associated PA; and (2) at least 1 first T helper cell group which is CC foreign to the animal. Analogues of human PSM, human Her2 and CC human/murine FGF8b comprising a substantial part of all known and CC predicted CMV and B-cell epitopes of the respective PA and including at CC least one foreign T helper epitope are also claimed. The method is used CC to treat prostate, prostate/breast or breast cancer when the PA is human CC PSM, FGF8b and Her2, respectively. Note: This sequence was constructed CC from the wild type human PSM (AAY92619), which appears on pages 184-187 CC of the specification

XX Sequence 750 AA;

SQ

Query Match 62.9%; Score 117; DB 3; Length 750;
Best Local Similarity 95.7%; Pred. No. 3e-08;
Matches 22; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 FNNFTVSEWLRLPKVSSASHLLEG 23
DB 32 FNNFTVSEWLRLPKVSSASHLLET 54

RESULT 13

ID AAY92636 standard; protein; 750 AA.

XX XX

AC AAY92636;

XX XX

DT 10-AUG-2000 (first entry)

XX DE Mutant human prostate specific membrane antigen construct, hPSM1.5.

XX KW Prostate specific membrane antigen; immunogenized construct; mutant;
KW vaccination; cytotoxic T-lymphocyte immunity; breast cancer;
KW prostate cancer; cell-associated peptide antigen; foreign epitope.

XX OS Homo sapiens.

XX Synthetic.

FH Key Peptide 24..38

FT FT /label= P2
FT /note= "foreign epitope"

FT Peptide 301..321

FT FT /label= P30
FT /note= "foreign epitope"

XX WO2000020027-A2.

PN PR 13-APR-2000.

XX PR 05-OCT-1999; 99WO-DK000525.

XX PR 05-OCT-1998; 9BDK-00001261.

XX PR 20-OCT-1998; 9BSN-0105011P.

XX PD 13-APR-2000.

XX PF 05-OCT-1999; 99WO-DK000525.

XX PR 05-OCT-1998; 9BDK-00001261.

XX PR 20-OCT-1998; 9BSN-0105011P.

XX PA (MBI-) M & E BIOTECH AS.

XX PI Steinnaa L, Mouritsen S, Nielsen KG,

PI Gautam A, Birk P, Karlsson G;

XX DR XX

XX WPI; 2000-349917/30.

XX PT Inducing immune responses to weakly immunogenic, tumor associated peptide
PR antigens for the treatment of breast and prostate cancer.

XX PS XX

CC AAY92627-49 are mutant immunogenized human prostate specific membrane CC antigen (PSM) constructs, which contain foreign epitopes (P2 and/or P30). CC The immunogenic analogues of PSM can be used in the claimed method as an CC adjuvaccine to induce a CTL response. Subdominant CTL epitopes, antibody CC binding regions and cysteine residues involved in disulfide bonds are CC preserved in the immunogenized forms. The method is used for inducing CC immune responses against weakly immunogenic cell-associated peptide CC antigens (PA) such as those associated with cancers (self-proteins), e.g.

antigen (PSM) constructs, which contain foreign epitopes (P2 and/or P30). The immunogenic analogues of PSM can be used in the claimed method as an autovaccine to induce a CTL response. Subdominant CTL epitopes, antibody binding regions and cysteine residues involved in disulfide bonds are preserved in the immunogenized forms. The method is used for inducing immune responses against weakly immunogenic cell-associated peptides (PA) such as those associated with cancers (self-proteins), e.g. human prostate specific membrane antigen (PSM), heregulin 2 (Her2) and/or fibroblast growth factor 8b (FGF8b). The method comprises effecting simultaneous presentation by antigen producing cells (APCs) of the animals immune system of: (1) at least 1 CTL (cytotoxic T-lymphocyte) group derived from the PA and/or least 1 B-cell group derived from the cell-associated PA; and (2) at least 1 first T helper cell group which is foreign to the animal. Analogues of human PSM, human Her2 and human/murine FGF8b comprising a substantial part of all known and predicted CTL and B-cell epitopes of the respective PA and including at least one foreign T helper epitope are also claimed. The method is used to treat prostate, prostate/breast or breast cancer when the PA is human PSM, FGFB and Her2, respectively. Note: This sequence was constructed from the wild type human PSM (AYA92619), which appears on pages 184-187 of the specification.

Sequence 750 AA;

Query Match	62.9%	Score 117;	DB 3;	Length 750;
Best Local Similarity	75.8%	Pred. No. 3e-08;	Indels 5;	Gaps 1;
Matches 25;	Conservative	1;	Mismatches 5;	

Qy 1 FNNFTVFWLRVPKVSASHLEGPSPLHWSYGLRP 33
Db 301 FNNFTVFWLRVPKVSASHLE--SLKVPTNVGP 331

RESULT 14
AYA84423 standard; protein; 188 AA.
XX AC AAY84423;
XX DT 25-JUL-2000 (first entry)

An osteoprotegerin ligand/tetanus toxoid P30 epitope fusion protein.
DE An osteoprotegerin ligand; OPG; osteoprotegerin; osteoclastogenesis;
KW tumour necrosis factor receptor; type II transmembrane protein;
KW osteoclast differentiation; CSF-1; osteoclast activator; immune response;
KW osteoporosis; bone resorption.
XX OS Synthetic.
OS Clostridium tetani.
OS Mus musculus.
XX Key Location/Qualifiers
Peptide 1..14 /note= "His tag"
FT Protein 15..112 /note= "residues 158-255 of murine OPG"
Peptide 113..133 /note= "tetanus toxoid P30 epitope"
Protein 134..188 /note= "residues 262-316 of murine OPG"
XX PN WO200115807-A1.
XX PD 23-MAR-2000.
XX PF 13-SEP-1999; 99WO-DK000481.

XX PR 15-SEP-1998; 98DK-00001164.
PR 02-OCT-1998; 98US-010286P.
XX PA (MEBI-) M & E BIOTECH AS.
PA (KLYS/) KLYSNER S.
PA (NIEL/) NIELSEN F S.

PI Halkier T, Haaning J;
XX WP1; 2000-27144/23.
DR N-PSDB; AAZ99970.
XX PT In vivo down-regulation of osteoprotegerin ligand (OPGL) activity used to treat, prevent and ameliorate osteoporosis.
XX PT Example; Page 94-95; 110PP; English.
PS The present sequence represents fusion protein of murine osteoprotegerin ligand (OPGL) and tetanus toxoid P30 epitope. Osteoprotegerin is a secreted member of the tumour necrosis factor receptor family, which blocks osteoclastogenesis in a dose dependent manner. The OPG protein is synthesised as a type II transmembrane protein. The murine and human OPG polypeptides are 87% homologous. OPG is a potent osteoclast differentiation factor when combined with CSF-1. It is not capable of inducing osteoclast differentiation in the absence of CSF-1. OPG is also an activator of mature osteoclasts. The specification describes a method for the in vivo down-regulation of OPGL activity in an animal. The method comprises using at least one OPGL polypeptide or subsequence, and/or at least one OPGI analogue to induce an immune response in the animal. The method and OPGI polypeptide are useful for treating, preventing and ameliorating osteoporosis or other diseases or conditions characterised by excessive bone resorption.

XX Sequence 188 AA;
SQ Score 116; DB 3; Length 188;
Best Local Similarity 82.1%; Pred. No. 8.6e-09;
Matches 23; Conservative 1; Mismatches 0; Indels 4; Gaps 1;

Qy 1 FNNFTVFWLRVPKVSASHLEGPSPLHWS 28
Db 113 FNNFTVFWLRVPKVSASHLE---NWS 136

RESULT 15
AA030458 ID AA030458 standard; protein; 285 AA.
XX AC AAO30458;
XX DT 22-SEP-2003 (first entry)
XX DE HILS-P2-P30-HILS (HILS:35) fusion construct Protein.
XX KW Multimeric protein; interleukin 5; IL5; TNFaLpha; inflammatory disease; tumour necrosis factor alpha; gene therapy; arthritis; interleukin 5; IL5; epitope; human; tetanus toxoid; chimeric.
XX Homo sapiens.
OS Unidentified.
OS Chimeric.
XX Key Location/Qualifiers
Peptide 1..19 /note= "Human IL5 leader peptide"
FT Peptide 20..285 /note= "Mature hIL5.35 protein"
FT Protein XX PN WO2003042244-A2.
XX PD 22-MAY-2003.
XX PP 15-NOV-2002; 2002WO-DK000764.
XX PR 16-NOV-2001; 2001DK-00001702.
PR 16-NOV-2001; 2001US-0331575P.
XX PA (PHAR) PHARMA XAS.
PA (KLYS/) KLYSNER S.
PA (NIEL/) NIELSEN F S.

PA (BRATT T.) BRATT T.
 PA (VOLD) VOLDBORG B.
 PA (MOURITSEN S.) MOURITSEN S.
 XX
 PI Klysner S., Nielsen FS, Bratt T, Voldborg B, Mouritzen S;
 XX
 DR WPI; 2003-449555/42.
 DR N-PSDD; AAI61294.
 XX
 BT New immunogenic analogue of a polymeric protein, useful for preparing a
 PR composition for treating inflammatory diseases e.g. arthritis.
 XX
 PS Claim 20; Page 112-113; 196pp; English.
 XX
 CC The invention relates to immunogenic analogues of multimeric proteins
 CC such as immunogenic variants of interleukin 5 (IL5) and tumour necrosis
 CC factor alpha (TNF_α, TNFalpha) and methods for production of immunogenic
 CC analogues. The immunogenic analogue is useful for preparing a composition
 CC for treating inflammatory diseases, e.g., arthritis. It is also used in
 CC gene therapy. The present sequence is a fusion construct which comprises
 CC 2 human interleukin 5 (IL5) monomers joined by tetanus toxoid epitopes
 CC P30 and P2. This sequence is used to illustrate the method of the
 CC invention
 XX
 Sequence 285 AA:

SQ Query Match 62.4%; Score 116; DB 6; Length 285;
 Best Local Similarity 91.7%; Pred. No. 1.4e-08;
 Matches 22; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 FNNFTVSEFWLRVPKVASHLEGPS 24
 Db 150 FNNFTVSEFWLRVPKVASHLEGPS 173

Search completed: March 10, 2004, 09:12:10
 Job time : 52.1984 secs

GenCore version 5.1.6
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CM protein - protein search, using sw model

Run on: March 10, 2004, 08:58:54 : Search time 10.5837 Seconds
 (without alignment)
 309.015 Million cell updates/sec

Title: US-09-848-834A-10
 Perfect score: 186
 Sequence: 1 PNNFTVSEWLRYVKVASHLEGPSLHWSYGLRPX 34

Scoring table: BLOSUM62
 Gapop 10.0 , Gapext 0.5

Searched: 283356 seqs, 96191526 residues
 Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0
 Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
 Maximum Match 100%
 Listing first 45 summaries

Database : PIR 78:
 1: pir1:
 2: pir2:
 3: pir3:
 4: pir4:
 *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

% Query Score Match Length DB ID Description

Result No.	Score	Query Match	Length	DB ID	Description
1	112	60.2	1315	1 BTCLTN	tentoxilysin (EC 3
2	61.5	33.3	1268	2 S33411	botulinum neurotoxin
3	61.5	33.1	92	1 RHRIG	gonadoliberin prec
4	61	32.8	366	2 S48110	neurotoxin type F
5	61	32.8	369	2 S48109	neurotoxin type F
6	61	32.8	1274	2 I40813	neurotoxin type F
7	61	32.8	1297	2 S39791	neurotoxin - Clost
8	59	31.7	1296	1 BTCLAB	bontoxilysin (EC 3
9	58	31.2	1291	1 A48940	bontoxilysin (EC 3
10	58	31.2	1291	2 I40631	non-proteolytic bo
11	57.5	30.9	90	1 RHMIG	gonadoliberin prec
12	57	30.6	502	2 T36589	probable transmemb
13	56.5	30.4	367	2 S48106	neurotoxin type F
14	56.5	30.4	1251	2 JH0256	botulinum neurotox
15	56.5	30.4	1252	2 S21178	botulinum neurotox
16	56	30.1	812	2 I01618	hypothetical prote
17	56	30.1	1296	2 I40635	botulinum neurotox
18	54.5	29.3	92	1 RHHUG	gonadoliberin prec
19	54.5	29.3	1285	2 S70582	botulinum neurotox
20	54.5	29.3	1291	2 S46431	botulinum neurotox
21	54.5	29.3	1291	2 A49777	gonadoliberin prec
22	54	29.0	67	2 I78541	probable alpha-amy
23	53.5	28.8	469	2 B37837	DNA-directed DNA P
24	53.5	28.8	3122	2 T17202	probable maturase
25	53	28.5	519	2 S78195	gonadoliberin - pi
26	52	28.0	10	1 RHPGG	gonadoliberin - gh
27	52	28.0	89	2 I51423	nucleocapsid prote
28	52	28.0	449	2 S23156	nucleocapsid prote

nonstructural prot
 nonstructural prot
 medaka-type Sonoda
 cytochrome-c Oxida
 probable maturase
 protein kinase (EC
 probable myb-like
 aconitase - Aquife
 hypothetical prote
 toxin, nontoxic co
 botulinum neurotox
 alpha Galactosytr
 hypothetical prote
 hypothetical prote
 sphaeroidine monoox
 ATP-binding casset

RESULT 1

BTCLTN tentoxilysin (EC 3.4.24.68) precursor - Clostridium tetani

N; Alternative names: tetanus neurotoxin

C; Species: Clostridium tetani

C; Sequence: 31-Mar-1988 #text change 31-Mar-1988 #text change 31-Jun-2002

C; Accession: A25659; A25757; S09348; S09364

R; Eisel, U.; Jarusch, W.; Goretzki, K.; Henschen, A.; Engels, J.; Waller, U.; Hudel, BMBJ J., 5, 2495-2502, 1986

A; Title: Tetanus toxin: primary structure, expression in E. coli, and homology with bovine insulin

A; Reference number: A25689; MUID:87053814; PMID:3536478

A; Accession: A25757; MUID:3774547

A; Residues: 1-1315 <E1S>

A; Cross-references: CB:X04436; NID:940769; PIDN:CAA28033.1; PID:940770

A; Cross-references: N.P.; Lyness, V.A.; Fairweather, N.P.; Lyness, V.A.; Pickard, D.J.; Allen, G.; Thomson, R.O.

R; Fairweather, N.P.; Lyness, V.A.; Nucleic Acids Res., 14, 7809-7811, 1986

J; Bacteriol., 165, 21-27, 1986

A; Title: Cloning, nucleotide sequencing, and expression of tetanus toxin fragment C in A; Reference number: A25194; MUID:86085672; PMID:3510187

A; Accession: A25194

A; Molecule type: DNA

A; Residues: 1-1315 <FA1>

A; Cross-references: GB:XM06214; NID:940773; PIDN:CAA29564.1; PID:940774

A; Molecule type: DNA

A; Residues: 1-1315 <FA1>

A; Cross-references: GB:X06214; NID:940773; PIDN:CAA29564.1; PID:940774

A; Experimental source: Strain CN391

R; Fairweather, N.P.; Lyness, V.A.; Nucleic Acids Res., 14, 7809-7811, 1986

J; Bacteriol., 165, 21-27, 1986

A; Title: The complete nucleotide sequence of tetanus toxin.

A; Reference number: A25757; MUID:3774547

A; Accession: A25757

A; Cross-references: GB:M12739; NID:914920; PIDN:AAA23282.1; PID:914921

A; Molecule type: DNA

A; Residues: 1-1315 <FA2>

A; Cross-references: B25194

A; Molecule type: Protein

A; Residues: B05-894 <FA3>

R; Matsuda, M.; Lei, D.L.; Sugimoto, N.; Ozutsumi, K.; Okabe, T.

Infect. Immun., 57, 3588-3591, 1989

J; Immunol., 142, 394-402, 1989

A; Title: Isolation, purification, and characterization of fragment B, the NH-2-termina

A; Reference number: A60759; MUID:9003436; PMID:12478376

A; Accession: A60759

A; Molecule type: Protein

A; Residues: 461-475 <MAT>

R; Demotz, S.; Lanzavecchia, L.; Eisel, U.; Niemann, H.; Widmann, C.; Corradini, G.

A; Title: Delineation of several DR-restricted tetanus toxin T cell epitopes.

A; Reference number: JS0098; MUID:8909318; PMID:2463305

R; Schiavo, G.; Benfenati, F.; Poulian, B.; Rossetto, O.; de Laureto, P.P.; DasGupta, B.

Nature, 335, 832-835, 1992

A; Title: Tetanus and botulinum-B neurotransmitter release by protein

A; Reference number: S27125; MUID:93063293; PMID:1331807

A; Contents: annotation

Ride Filippis, V.; Vangelista, L.; Schiavo, G.; Tonello, F.; Montecucco, C.
Eur. J. Biochem. 229, 61-69, 1995
A;Title: Structural studies on the zinc-endopeptidase light chain of tetanus neurotoxin.

A;Reference number: 663348; PMID:7744050

A;Accession: S69348
A;Molecule type: Protein

A;Residues: 2-31 <DBP>

C;Comment: The source of this protein was an extrachromosomal plasmid.

C;Comment: The precursor is cleaved by endogenous proteinase activity to form light (fragment B) dual chains. They are not toxic when separated.

C;Comment: Fragment B forms ion channels in a lipid bilayer. Fragment C binds to ganglionic synapses; it is identical to hypothalamic luteinizing hormone.

C;Comment: This potent neurotoxin binds to peripheral neuronal synapses; it is identical to hypothalamic luteinizing hormone.

C;Function: Presynaptic neurons. It inhibits neurotransmitter release by proteolytic cleavage of synaptic vesicle protein.

C;Description: Blocks neurotoxicosis via hydrolysis of a Glu-Phe peptide bond in synaptosomal membrane protein.

C;Superfamily: tetanus toxin

C;Keywords: hydrolase; metalloprotease; neurotoxin; transmembrane protein; zinc

F;461-1315/Product: tentoxysin light chain (fragment A) #status predicted <URL>

F;461-1315/Product: tentoxysin heavy chain (fragment B-C) #status experimental <TH>

F;865-1315/Domain: channel forming (fragment B) #status predicted <TXB>

F;233-237/Domain: ganglioside binding (fragment C) #status predicted <TXC>

F;234/Active site: zinc (His) #status predicted

F;234/Active site: Glu #status predicted

Query Match 60.2%; Score 112; DB 1; Length 1315;

Best Local Similarity 100.0%; Pred. No. 2e-07;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 FNNFTYFWLRYPKVSAHLE 21

Db 947 FNNFTYFWLRYPKVSAHLE 967

RESULT 2

botulinum neurotoxin type F - Clostridium baratii

C;Species: Clostridium baratii

C;Date: 13-Jan-1995 #sequence_revision 13-Jan-1995 #text_change 16-Jul-1999

C;Accession: S33411; S31860

R;Thompson, D.B.; Huston, R.A.; East, A.K.; Allaway, D.; Collins, M.D.; Richardson, P.T.

PEMS Microbiol. Lett. 108, 175-182, 1993

A;Title: Nucleotide sequence of the gene coding for Clostridium baratii type F neurotoxin

A;Reference number: S33411; MUID:8486245

A;Accession: S33411

A;Status: Preliminary

A;Molecule type: DNA

A;Residues: 1-1268 <THO>

A;Cross-references: EMBL:X682622; PID:949138; PID:CAA48329.1; PMID:949139

C;Superfamily: tetanus toxin

C;Keywords: neurotoxin

Query Match 33.3%; Score 62; DB 2; Length 1268;

Best Local Similarity 64.3%; Pred. No. 2.2%;
Matches 9; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Qy 1 FNNFTYFWLRYPK 14

Db 922 YQNFSYFWWRIPK 935

RESULT 3

JHRTG

Gonadotropin precursor - rat

N;Alternative names: gonadotropin-associated protein (GAP); gonadotropin releasing hormone

C;Contains: Gonadotropin precursor - rat

C;Species: Rattus norvegicus (Norway rat)

C;Date: 31-Mar-1988 #sequence_revision 31-Mar-1988 #text_change 18-Jun-1999

C;Accession: A40147; B26173; A48410

R;Bond, C.T.; Hayfllick, J.S.; Seeburg, P.H.; Adelman, J.P.

Mol. Endocrinol. 3, 1257-1262, 1989

A;Title: The rat Gonadotropin-releasing hormone: SH locus: structure and hypothalamic ex-

A;Reference number: A40147; MUID:83384661; PMID:2476669

A;Accession: A40147

A;Molecule type: DNA

A;Residues: 1-92 <DBN>

A;Cross-references: GB:M31670; PID:920447; PID:AAA41264.1; PMID:920448

R;Adelman, J.P.; Mason, A.J.; Hayfllick, J.S.; Seeburg, P.H.

Proc. Natl. Acad. Sci. U.S.A. 83, 179-183, 1986

A;Title: Isolation of the gene and hypothalamic cDNA for the common precursor of gonado-

A;Reference number: A94090; MUID:86094338; PMID:2867548

A;Accession: B26173

A;Molecule type: mRNA

A;Residues: 1-92 <ADE>

A;Cross-references: GB:M12579; PID:920445; PID:920446

R;Leboeuf, B.; Blalock, J.E.

Proc. Natl. Acad. Sci. U.S.A. 83, 179-183, 1986

A;Title: Isolation of the gene and hypothalamic cDNA for the common precursor of gonado-

A;Reference number: A94090; MUID:86094338; PMID:2867548

A;Accession: A48410

A;Status: preliminary

A;Molecule type: mRNA

A;Residues: 1-92 <NAI>

A;Cross-references: GB:S50870; PID:9262060

A;Experimental source: thymus

A;Title: Thymocytes express a mRNA that is identical to hypothalamic luteinizing hor-

A;Reference number: A93105480; PMID:1468115

A;Accession: A48410

A;Function:

C;Genetics:

A;Interventions:

C;Keywords:

A;Description: stimulates pituitary secretion of lutropin and follitropin

A;Note: gonadotropin-associated protein may have prolactin release inhibiting activity

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

C;Superfamily: Gonadotropin

C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglutamic acid

RESULT 5

S48109
neurotoxin type F - Clostridium botulinum (fragment)
C;Species: Clostridium botulinum
C;Accession: 12-Feb-1998 #sequence_revision 20-Feb-1998 #text_change 16-Jul-1999
R;Campbell, K.D.; Collins, M.D.; East, A.K.
J;Clin. Microbiol. 31, 2255-2262, 1993
A;Title: Gene probes for identification of the botulinum neurotoxin gene and specific id
A;Reference number: S48103; MUID:940133; PMID:8408542
A;Status: preliminary; nucleic acid sequence not shown; translation not shown
A;Residues: 1-369 <CAM>
A;Cross-references: EMBL:X70820; NID:9407790; PIDN:CAA50151.1; PMID:9407791
A;Note: the nucleotide sequence was submitted to the EMBL Data Library, January 1993
C;Superfamily: tetanus toxin

Query Match 32.8%; Score 61; DB 2; Length 369;
Best Local Similarity 57.1%; Pred. No. 0.77%;
Matches 8; Conservative 5; Mismatches 1; Indels 0; Gaps 0;
Qy 1 FNNFTVSFWLRVPK 14
Db 297 YQNFSISFWVRIPK 310

RESULT 6
140813
A;Status: preliminary; translated from GB/EMBL/DDBJ
A;Molecule type: DNA
A;Residues: 1-174 <SBS>
A;Cross-references: GB:MR92906; NID:9144866; PIDN:AAA23263.1; PMID:9144867
R;Campbell, K.D.; Collins, M.D.; East, A.K.
J;Clin. Microbiol. 31, 2255-2262, 1993
A;Title: Gene probes for identification of the botulinum neurotoxin gene and specific id
A;Reference number: S48103; MUID:940133; PMID:8408542
A;Accession: S48108
A;Status: preliminary; translation not shown
A;Molecule type: DNA
A;Residues: 634-1002 <CAM>
A;Cross-references: EMBL:X70816; NID:9407788; PIDN:CAA50147.1; PMID:9407789
C;Superfamily: tetanus toxin
C;Keywords: neurotoxin

Query Match 32.8%; Score 61; DB 2; Length 1274;
Best Local Similarity 57.1%; Pred. No. 3%;
Matches 8; Conservative 5; Mismatches 1; Indels 0; Gaps 0;
Qy 1 FNNFTVSFWLRVK 14
Db 930 YQNFSISFWVRIPK 943

RESULT 7
S39791
neurotoxin - Clostridium botulinum
C;Species: Clostridium botulinum
C;Accession: 07-Oct-1994 #sequence_revision 01-Dec-1995 #text_change 16-Jul-1999
R;Campbell, K.; Collins, M.D.; East, A.K.
Biochim. Biophys. Acta 1216, 487-491, 1993
A;Title: Nucleotide sequence of the gene coding for Clostridium botulinum (Clostridium a
A;Reference number: S39791; MUID:94092745; PMID:82288233
A;Accession: S39791

A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-1297 <CAM>
A;Cross-references: EMBL:X74162; NID:9441275; PIDN:CAA52275.1; PMID:9441276
C;Superfamily: tetanus toxin
C;Keywords: neurotoxin
Query Match 32.8%; Score 61; DB 2; Length 1297;
Best Local Similarity 38.1%; Pred. No. 3.1%;
Matches 8; Conservative 9; Mismatches 4; Indels 0; Gaps 0;
Qy 1 FNNFTVSFWLRVPKSASHLE 21
Db 930 FDNFSINFWRTPKNNNDIQ 950

RESULT 8
ENCLAB
bontoxilysin (EC 3.4.24.69) A precursor - Clostridium botulinum
N;Alternative names: botulinum neurotoxin type A
C;Species: Clostridium botulinum
C;Date: 31-Mar-1993 #sequence_revision 31-Mar-1993 #text_change 18-Jun-1999
C;Accession: A35294; A60725; A60726
R;Binz, T.; Kurazono, H.; Willse, M.; Fravert, J.; Niemann, H.
J. Biol. Chem. 265, 9153-9158, 1990
A;Title: The complete sequence of botulinum neurotoxin type A and comparison with other
A;Reference number: A35294; MUID:90164400; PMID:2160960
A;Accession: A35294
A;Molecule type: DNA
A;Residues: 1-1296 <BIN>
A;Cross-references: GB:M30196; NID:9144864; PIDN:AAA23262.1; PMID:9144865
A;Experimental source: strain 62A, subtype A
R;Thompson, D.E.; Brehm, J.K.; Cultrami, J.D.; Swinfield, T.J.; Shone, C.C.; Atkinson,
Eur. J. Biochem. 189, 73-81, 1990
A;Title: The complete amino acid sequence of the Clostridium botulinum type A neurotoxin
A;Reference number: S09492; MUID:90235864; PMID:2185020
A;Accession: S09492
A;Molecule type: DNA
A;Residues: 1,'Q',3-26,'V',28-1296 <THO>
A;Cross-references: EMBL:Z2066; NID:940381; PIDN:CNA36289.1; PMID:940382
A;Experimental source: NCTC 2916
R;Fujita, R.; Fujinaga, Y.; Inoue, K.; Nakajima, H.; Kumon, H.; Oguma, K.
FEBS Lett. 376, 41-44, 1995
A;Title: Molecular characterization of two forms of nontoxic-nonhemagglutinin component
A;Reference number: S67988; MUID:96096783; PMID:8521962
A;Accession: S67988
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-12 <SFUJ>
A;Cross-references: EMBL:D67030; DDBJ:D50421; NID:92160224
R;Batley, M.J.; Somers, E.; DasGupta, B.R.
Biochim. Biophys. Res. Commun. 162, 1388-1395, 1989
A;Title: Characterization of botulinum type A neurotoxin gene: delineation of the N-terminal
A;Reference number: A33401; MUID:89305959; PMID:2669749
A;Accession: A33401
A;Molecule type: DNA
A;Residues: 1-35 <BT>
A;Cross-references: GB:M27892; NID:9144880; PIDN:AAA23263.1; PMID:95511776
R;Gimenez, J.A.; DasGupta, B.R.
J. Protein Chem. 12, 351-363, 1993
A;Title: Botulinum type A neurotoxin digested with pepsin yields 132, 97, 72, 45, 42,
A;Reference number: A53884; MUID:8397793
A;Accession: A53884
A;Status: preliminary
A;Molecule type: protein
A;Residues: 867-880; 114-1817,'Y',1219 <GIM>
A;Experimental source: strain Hall
A;Note: sequence extracted from NCBI backbone (NCBIP:139159); sequence modified after
R;DasGupta, B.R.; Dekleva, M.L.
Biochim. Biophys. Acta 1216, 661-664, 1990
A;Reference number: A60025; MUID:91120847; PMID:2126206
A;Accession: A60025

A;Title: Minimal essential domains specifying toxicity of the light chains of tetanus type A
A;Molecule type: protein
A;Residues: 2-6;4453-'X';455-457 <Das1>
A;Reference number: A42871; MUID:92340309; PMID:1634516
A;Accession number: A2871
A;Status: nucleic acid sequence not shown
A;Molecule type: mRNA
A;Residues: 1-313; 'S'; 315-451 <KUR>
A;Note: Sequence extracted from NCBI backbone (NCBIP:109365)
R;DasGupta, B.R.; Datta, A.
Biochim Biophys Acta 70, 811-817, 1988
A;Title: Botulinum neurotoxin type B (strain 657): partial sequence and similarity with
A;Reference number: S07155; MUID:89000387; PMID:139097
A;Accession: S0562
A;Molecule type: protein
A;Residues: 2-29; 'M'; 31-45 <Das>>
A;Molecule type: protein
A;Residues: 442-463; 'R'; 465-467 <DA2>
R;Schmidt, J.J.; Sathyamoorthy, V.; DasGupta, B.R.
Arch. Biochem. Biophys. 238, 544-548, 1985
A;Title: Partial amino acid sequences of botulinum neurotoxins types B and E.
A;Reference number: S07128; MUID:85197963; PMID:3888113
A;Accession: S07128
A;Molecule type: protein
A;Residues: 2-17 <SCH1>
A;Molecule type: protein
A;Accession: S08573
A;Status: Preliminary
A;Molecule type: protein
A;Residues: 2-17 <SCH2>
A;Accession: S08574
A;Status: Preliminary
A;Molecule type: protein
A;Accession: 442-459 <SCH3>
R;Schiavo, G.; Benfenati, F.; Poulat, B.; Rossetto, O.; de Laureto, P.P.; DasGupta, B.
Nature 359, 832-835, 1992
A;Title: Tetanus and botulinum-B neurotoxins block neurotransmitter release by proteolytic
A;Contents: annotation
C;Comment: Botulinum neurotoxins inhibit neurotransmitter release from cholinergic syna
C;Genetics:
A;Gene: atx; batA
C;Function:
A;Molecule type: protein
A;Residues: 2-47 <DS2>
A;Description: catalyzes hydrolysis of an Asn-Arg peptide bond in synaptosomal-associate
C;Superfamily: tetanus toxin
C;Keywords: metalloproteinase; neurotoxin; transmembrane pro
F;445-1296/Product: bontoxilysin A light chain #status experimental <LIGHT>
F;445-1297/Product: bontoxilysin A heavy chain #status experimental <HHV>
F;223-227/Binding site: zinc [His] #status predicted
F;224/Active site: Glu #status predicted
Query Match Score 59; DB 1; Length 1296;
Best Local Similarity 57.1%; Pred. No. 5.9;
Matches 8; Conservative 4; Mismatches 2; Indels 0; Gaps 0;
Oy 1 FNNFTYSEFWLWPK 14
Db 938 YENFSTSFWIRPK 951
RESULT 9
A48940
bontoxilysin (EC 3.4.24.69) B precursor - Clostridium botulinum
N;Alternative names: botulinum neurotoxin type B (BNT/B)
C;Species: Clostridium botulinum
C;Date: 19-Dec-1993 #sequence revision 18-Nov-1993 #text change 18-Jun-1999
C;Accession: A48940; S48105; S21575; A42871; S07155; S08562; S07128; S08574
R;Whelan, S.M.; Elmore, M.J.; Bodsworth, N.J.; Brim, J.K.; Atkinson, T.; Minton, N.P.
APPL Environ. Microbiol. 58, 2343-2354, 1992
A;Title: Molecular cloning of the Clostridium botulinum structural gene encoding the typ
A;Reference number: A48940; MUID:92384550; PMID:1514783
A;Accession: A48940
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-1231 <THE>
A;Cross-references: GB:MB1186; NID:9144734; PID:9144735
A;Experimental source: type B; Danish
A;Note: sequence extracted from NCBI backbone (NCBIN:112080; NCBIPI:112081); this publica
R;Campbell, K.D.; Collins, M.D.; East, A.K.
J Clin. Microbiol. 31, 2255-2262, 1993
A;Title: Gene probes for identification of the botulinum neurotoxin gene and specific id
A;Reference number: S48103; MUID:94013372; PMID:8408542
A;Accession: S48105
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 634-994 <CRM>
A;Cross-references: EMBL:X70817; NID:9407782; PID:9407783
A;Experimental source: proteolytic type B; strain NCTC 7273
R;Szabo, E.A.; Pemberton, J.M.; Desmarchier, F.M.
submitted to the EMBL Data Library, April 1992
A;Description: Partial amino acid sequence of botulinum neurotoxin type B and comparisio
A;Reference number: S21575
A;Accession: S21575
A;Molecule type: DNA
A;Residues: 36-217; 'G'; 219-224; 'S'; 226-246 <SSZ>
A;Cross-references: EMBL:Z11934; NID:940383; PID:940384
R;Kurazono, H.; Mochida, S.; Bintz, T.; Eisell, U.; Quanz, M.; Grebenstein, O.; Wernars, R.
T;Dissertation
RESULT 10
I40331
non-proteolytic botulinum neurotoxin type B precursor - Clostridium botulinum
C;Species: Clostridium botulinum
C;Date: 12-Aug-1996 #sequence 14-37; P-Prod. No. 8.1; Length 1291;
Best Local Similarity 64.3%; P-Mismatches 0; Gaps 0;
Matches 9; Conservative 4; Mismatches 1; Indels 0;
R;Hutson, R.A.; Collins, M.D.; East, A.K.; Thompson, D.E.
Curr. Microbiol. 28, 101-110, 1994
A;Title: Nucleotide sequence of the gene coding for non-proteolytic Clostridium botulinum
A;Reference number: I40631; MUID:94122659; PMID:7764370
A;Accession: I40631
A;Molecule type: DNA
A;Cross-references: GB:EMBL/DBJ

A;Accession: JH0256
A;Status: nucleic acid sequence not shown
A;Molecule type: DNA
A;Residues: 1-27, 'E', 29-1251 <POU>
A;Cross-references: EMBL:X3180; NID:940379
A;Experimental source: strains ATCC 43181 and ATCC 43755
R;Fujii, N.; Kimura, K.; Yashiki, T.; Indoh, T.; Murakami, T.; Tsuzuki, K.; Yokosawa, N.
J. Gen. Microbiol. 137, 519-525, 1991
A;Title: Cloning of a DNA fragment encoding the 5'-terminus of the botulinum type E toxin
A;Reference number: S16145; MUID:91237316; PMID:2033376
A;Accession: S16145
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-229, 'M', 231-252 <POU>
A;Cross-references: EMBL:X3180; NID:940407; PIDN:CAA37321.1; PID:940408
A;Experimental source: strain B6340
C;Comment: The clostridial neurotoxins are toxins that inhibit neurotransmitter release
C;Comment: The heavy chain mediates the binding of toxin to cell receptors while the light chain mediates the binding of toxin to cell receptors while the ligand binds to the receptor
C;Superfamily: tetanus toxin
C;Keywords: neurotoxin
P;422-422/Product: botulinum neurotoxin type E fragment encoding the 5'-terminus of the botulinum type E toxin
P;423-425/Product: botulinum neurotoxin type E fragment encoding the 5'-terminus of the botulinum type E toxin
P;412-426/Disulfide bonds: #status predicted
P;412-426/Disulfide bonds: #status predicted

A;Title: The complete sequence of botulinum neurotoxin type A and comparison with other neurotoxins
A;Reference number: A35294; MUID:90264400; PMID:2160960
A;Accession: B35294
A;Status: not compared with conceptual translation
A;Molecule type: DNA
A;Residues: 1-176, 'R', 178-252 <BIN>
A;Experimental source: strain Beluga
R;Gimenez, J.A.; DasGupta, B.R.
Biochim. Biophys. Acta 72, 213-217, 1990
A;Title: Botulinum neurotoxin type E fragment with endoproteinase Lys-C reveals the structure of the light chain
A;Reference number: A60027; MUID:90344918; PMID:2116911
A;Accession: A60027
A;Molecule type: Protein
A;Residues: 420-427 <GM>
A;Experimental source: strain Beluga
A;Note: this fragment was generated by proteolysis with Lys-C rather than with trypsin
C;Comment: The clostridial neurotoxins are highly potent protein toxins that inhibit the binding of toxin to cell receptors while the light chain mediates the binding of toxin to cell receptors while the heavy chain binds to the receptor
C;Superfamily: tetanus toxin
C;Keywords: neurotoxin
P;423-425/Product: botulinum neurotoxin type E light chain #status predicted <HEA>
P;412-426/Disulfide bonds: #status predicted
P;412-426/Disulfide bonds: #status predicted

Query Match Score 56.5; DB 2; Length 1252;
Best Local Similarity 22.4%; Pred. No. 13;
Matches 15; Conservative 8; Mismatches 7; Indels 37; Gaps 2;

Qy	1 FNNFTYSFWLRVP-----KVSASHLE-----	GP 23	
Db	912 YKNFTYSFWLRVPNYDNKIVVNNNEYTIINCMRDNNSSWIKVSLNHNELIWTIQDNSSINQ 971	Db	
Qy	24 SLHWSYG 30	Qy	24 SLHWSYG 30
Db	972 KLAFLNYG 978	Db	972 KLAFLNYG 978

Search completed: March 10, 2004, 09:16:46
Job time : 11.6425 secs

RESULT 15
S21178
Botulinum neurotoxin type E precursor - Clostridium botulinum
C;Species: Clostridium botulinum
C;Dte: 30-Sep-1993 #sequence revision 30-Sep-1993 #text change 15-Oct-1999
C;Cross-references: S21178; S48107; S51811
R;Whelan, S.M.; Elmore, M.J.; Bodsworth, N.J.; Atkinson, T.; Ninton, N.P.
Bur. J. Biochem. 204, 657-667, 1992
A;Title: The complete amino acid sequence of the Clostridium botulinum type-E neurotoxin
A;Reference number: S21178; MUID:92174922; PMID:1541280
A;Accession: S21178
A;Molecule type: DNA
A;Residues 1-1252 <WHE>
A;Cross-references: EMBL:X62683; NID:940397; PIDN:CAA44555.1; PID:940398
R;Campbell, K.D.; Collins, M.D.; East, A.K.
J. Clin. Microbiol. 31, 2255-2262, 1993
A;Title: Gene probes for identification of the botulinum neurotoxin gene and specific identification of the botulinum neurotoxin E derived from Clostridium botulinum type E
A;Reference number: JH0256; MUID:92181428; PMID:1543481
A;Accession: JH0257
A;Status: nucleic acid sequence not shown
A;Molecule type: DNA
A;Residues: 1-176, 'R', 178-197, 'C', 199-339, 'R', 341-772, 'I', 774-962, 'PE', 965-966, 'R', 968-1
A;Cross-references: EMBL:X62089; NID:940393; PIDN:CAA43996.1; PMID:91394
A;Experimental source: strain Beluga
R;Binz, T.; Kurzono, H.; Wille, M.; Frevert, J.; Wernars, K.; Niemann, H.
J. Biol. Chem. 265, 9153-9158, 1990

Scoring table:	BLOSUM62				
Gapop 10.0 , Gapext 0.5					
Searched:	141681 seqs, 52070155 residues				
Total number of hits satisfying chosen parameters:	1416881				
Minimum DB seq length:	0				
Maximum DB seq length:	2000000000				
Post-processing:	Minimum Match 0% Maximum Match 100% Listing first 45 summaries				
Database :	SwissProt_42;*				
Pred. No.	is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.				
SUMMARIES					
Result No.	Score	Query Match	Length	DB ID	Description
1	112	60 2	1314	1 TETX_CLOTE	P0458 clostridium
2	61	33 9	431	1 PURA_CLOSPN	Q84rm2 legionella
3	61.5	33 1	92	1 GONI_RAT	P0790 rattus norv
4	61	32 8	1274	1 BXF_CLOBO	P30996 clostridium
5	61	32 8	1296	1 BXG_CLOBO	Q63433 clostridium
6	59	31 7	1295	1 BXI_CLOBO	P10845 clostridium
7	58.5	31 5	1250	1 BXE_CLOBO	Q00496 clostridium
8	58	31 2	1290	1 BXB_CLOBO	P10446 clostridium
9	57.5	30 9	1051	1 GONI_MOUSE	P13522 mus musculus
10	57	30 6	1051	1 VP2_AHSV6	Q71024 african hor
11	56.5	30 4	90	1 GONI_RANCA	Q90163 ran catesbe
12	56.5	30 4	1250	1 BXE_CLOBO	P30995 clostridium
13	56	30 1	1295	1 BXZ_CLOBO	Q45694 clostridium
14	54.5	29 3	92	1 GONI_HUMAN	P10148 homo sapien
15	54.5	29 3	1290	1 BXC1_CLOBO	P18440 clostridium
16	54	29 0	67	1 GONI_MACMU	P55247 macaca mulatta
17	54	29 0	91	1 GONI_PIG	P49321 sus scrofa
18	53.5	28 8	760	1 AMY_CLOAB	P23671 clostridium
19	53.5	28 8	3122	1 DPOZ_MOUSE	Q61493 mus musculus
20	52	28 0	467	1 GONI_SHEEP	P20588 ovis aries
21	52	28 0	63	1 GONI_MESAU	Q93163 mesocricetus
22	52	28 0	89	1 GONI_XENLA	P45656 xenopus laevis
23	52	28 0	92	1 GONI_TUPGB	Q93335 tupala glis
24	52	28 0	449	1 VNNS_INSVN	Q08111 impatiens n
25	52	28 0	464	1 VNNS_TSIVW1	P26002 tomato spot
26	52	28 0	467	1 VNNS_TSIVW1	P26003 tomato spot
27	51.5	27 7	90	1 GON8 RANDY	Q9au2 rana dybowski
28	51.5	27 7	91	1 GON1_ORYLA	Q9dg8 o proponado
29	51.5	27 7	521	1 UBP3_HUMAN	Q9614 homo sapien
30	51	27 4	95	1 GON1_MORSA	Q73812 morone saxatilis
31	51	27 4	265	1 DDHC_RHOSU	Q8pg1 rhoovulum
32	51	27 4	292	1 C22_ORYSA	P29619 oryza sativa
33	51	27 4	1196	1 BXCN_CLOBO	P4081 clostridium

ALIGNMENTS

RESULT 1	TETX_CLOTE	STANDARD;	PRT;	1314 AA.
ID	TETX_CLOTE			
AC	P0958;			
DT	13-AUG-1987 (Rel. 05, Created)			
DT	13-AUG-1987 (Rel. 05, Last sequence update)			
DT	10-OCT-2003 (Rel. 42, Last annotation update)			
DE	Tetanus toxin Precursor (EC 3.4.24.68) ("Tetraoxlysin") [Contains: Tetanus toxin heavy chain (Tetanus toxin chain L); Tetanus toxin heavy chain (Tetanus toxin chain H); Tetanus toxin heavy chain (Tetanus toxin chain H).]			
GN	TETX OR CTPE60.			
OS	Clostridium tetani.			
OG	Plasmid PE88, and Plasmid 75 Kbp.			
OC	Bacteria; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;			
OX	NCBI_TaxID=1513;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	PLASMID=75 kbp;			
RX	MEDLINE=87053814; PubMed=3536478;			
RA	Eisele U., Jarausch W., Goretzki K., Henschchen A., Niemann H., Engels J.,			
RA	Weller U., Hudel M., Habermann E., Niemann H., Weller N.F., Fricke W.F., Liesegang H., Decker I., Herzerberg C., Martinez-Arias R., Merkl R., Henne A., Gottschalk G.;			
RT	"Tetanus toxin: primary structure, expression in E. coli, and homology with botulinum toxins.";			
RL	EMBO J. 12:2495-2502(1998).			
RN	[2]			
RP	SEQUENCE FROM N.A.			
RC	PLASMID=75 kbp;			
RX	MEDLINE=87040747; PubMed=3774547;			
RA	Fairweather N.F., Lyness V.A., Fairweather N.F., Lyness V.A.;			
RA	"The complete nucleotide sequence of tetanus toxin.";			
RT	RNA			
RL	Nucleic Acids Res. 14:7805-7812(1996).			
RN	[3]			
RP	SEQUENCE FROM N.A.			
RC	SEQUENCE FROM N.A.			
RX	MEDLINE=12457253; PubMed=12555129;			
RA	Brueggemann H., Baumer S., Liesegang H., Decker I., Herzerberg C., Gottschalk G.;			
RA	"Cloning, nucleotide sequencing, and expression of tetanus toxin fragment C in Escherichia coli.";			
RT	J. Bacteriol. 165:21-27(1986).			
RL	[4]			
RP	SEQUENCE OF 742-1314 FROM N.A.			
RC	PLASMID=75 kbp;			
RX	MEDLINE=86808562;			
RA	Fairweather N.F., Lyness V.A., Pickard D.J., Allen G., Thomson R.O., Kriegstein K., Henschchen A., Weller U., Habermann E.,			
RA	"PARTIAL SEQUENCE, AND DISULFIDE BONDS.			
RT	RNA			
RL	Proc. Natl. Acad. Sci. U.S.A. 100:1316-1321(2003).			
RN	[5]			
RP	SEQUENCE OF 742-1314 FROM N.A.			
RC	PLASMID=75 kbp;			
RX	MEDLINE=90201034;			
RA	Kriegstein K., Henschchen A., Weller U., Habermann E.,			
RA	"Arrangement of disulfide bridges and positions of sulphydryl groups in tetanus toxin.";			

Bur. J. Biochem. 188:39-45(1990).

[6] PARTIAL SEQUENCE; PubMed=1935979;
 RX Kriegstein K.G.; Henschel A.H.; Weller U.; Habermann E.;
 RA "Limited proteolysis of tetanus toxin. Relation to activity and
 RT identification of cleavage sites.";
 RL Bur. J. Biochem. 202:41-51(1991).
 RN RP IDENTIFICATION AS ZINC-PROTEASE;
 MEDLINE=93010948; PubMed=3396558;
 RA Schiavo G., Pouillain B., Rossetto O., Benfenati F., Tauc L.,
 RA Montecucco C.;
 RT "Tetanus toxin is a zinc protein and its inhibition of
 neurotransmitter release and protease activity depend on zinc.";
 RL EMBO J. 11:3577-3583(1992).
 RN RP IDENTIFICATION OF SUBSTRATE; PubMed=1331807;
 RX MEDLINE=93063293; PubMed=1331807;
 RA Schiavo G., Benfenati F., Pouillain B., Rossetto O., de Laureto P.P.,
 RA D'Agripa B.R., Montecucco C.;
 RT "Tetanus and botulinum-B neurotoxins block neurotransmitter release
 by proteolytic cleavage of synaptobrevin.";
 RA Nature 359:832-835(1992).
 RN RP X-RAY CRYSTALLOGRAPHY (2.7 ANGSTROMS) OF 874-1314.
 RX MEDLINE=97475217; PubMed=9334741;
 RA Umland T.C., Wingert L.M., Swaminathan S., Furey W.F., Schmidt J.J.,
 RA Sax M.;
 RT "Structure of the receptor binding fragment HC of tetanus
 RT neurotoxin.",
 RL Nat. Struct. Biol. 4:788-792(1997).
 CC |- FUNCTION: TETANUS TOXIN ACTS BY INHIBITING NEUROTRANSMITTER
 RELEASE. IT BINDS TO PERIPHERAL NEURONAL SYNAPSES, IS INTERNALIZED
 AND MOVES BY RETROGRADE TRANSPORT UP THE SPINAL CORD
 WHERE IT CAN MOVE BETWEEN POSTSYNAPTIC AND PRESYNAPTIC NEURONS. IT
 INHIBITS NEUROTRANSMITTER RELEASE BY ACTING AS A ZINC
 ENDOPROTEIDASE THAT CATALYZES THE HYDROLYSIS OF THE 76-GLN-1-PHE-77
 CC BOND OF SYNAPTOBREVIN-2.
 CC |- CATALYTIC ACTIVITY: Hydrolysis of 76-Gln-1-Phe-77 bond in
 CC synaptobrevin 2.
 CC |- COFACTOR: Binds 1 zinc ion per subunit (By similarity).
 CC |- SUBUNIT: THE PRECURSOR POLYPEPTIDE IS SUBSEQUENTLY CLEAVED TO
 CC YIELD SUBCHAINS L AND H. THESE REMAIN LINKED BY A DISULFIDE BRIDGE
 CC AND ARE NON-TOXIC AFTER SEPARATION.
 CC |- MISCELLANEOUS: THE C-TERMINAL OF THE HEAVY CHAIN BINDS TO
 CC GANGLIOSIDE RECEPTORS.
 CC |- SIMILARITY: Belongs to peptidase family M27.

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CC DR EMBL: X04436; CAA38033; 1; .
 CC DR EMBL: X05214; CAA32564; 1; .
 CC DR EMBL: AF28097; AAQ03749; 1; .
 CC DR EMBL: M12739; AAA3282; 1; .
 CC DR PIR: A25639; BTCLTN.
 CC DR PDB: 1AF9; 29-APR-98.
 CC DR PDB: 1ABD; 14-OCT-98.
 CC DR PDB: 1DDH; 27-MAR-00.
 CC DR PDB: 1DFQ; 24-MAR-00.
 CC DR PDB: 1DIW; 24-MAR-00.
 CC DR PDB: 1DLI; 24-MAR-00.
 CC DR PDB: 1FV3; 05-SEP-01.
 CC DR MEROPS: M27_001; .
 CC DR InterPro: IPR00898; ConA_like_lec_g1.
 CC DR InterPro: IPR00216; Kunitz_legume.
 CC DR InterPro: IPR006025; Pept_M_2n_BS.
 DR DR InterPro: IPR000355; Peptidase_M27.
 DR DR Pfam: PF01742; Peptidases_M27; 1.
 DR DR PRINTS: PR0070; BONTOXI_M27.
 DR DR ProDom: P000163; Bontoxin; 1.
 DR DR PROSITE; PS0042; ZINC_PROTOSIN; 1.
 KW KW Neurotoxin; Transmembrane_Hydrolease; Metalloprotease; Zinc; Plasmid;
 KW KW 3D-structure; Complete proteome.

FT FT INIT MET 0 0
 FT FT CHAIN 1 456
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 FT FT HELIX 873 882
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 FT FT STRAND 884 891
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 FT FT STRAND 923 925
 FT FT TURN 932 935
 FT FT STRAND 936 940
 FT FT HELIX 941 946
 FT FT STRAND 949 956
 FT FT TURN 956 962
 FT FT HELIX 968 970
 FT FT TURN 969 970
 FT FT STRAND 972 977
 FT FT STRAND 980 981
 FT FT HELIX 983 985
 FT FT STRAND 987 995
 FT FT TURN 996 997
 FT FT STRAND 998 1004
 FT FT TURN 1006 1007
 FT FT STRAND 1010 1016
 FT FT STRAND 1020 1020
 FT FT TURN 1021 1022
 FT FT STRAND 1031 1037
 FT FT TURN 1039 1040
 FT FT STRAND 1042 1047
 FT FT TURN 1048 1049
 FT FT STRAND 1050 1056
 FT FT TURN 1058 1059
 FT FT STRAND 1068 1074
 FT FT TURN 1079 1080
 FT FT STRAND 1082 1091
 FT FT HELIX 1097 1105
 FT FT TURN 1106 1107
 FT FT STRAND 1112 1112
 FT FT STRAND 1114 1114
 FT FT TURN 1116 1117
 FT FT STRAND 1120 1120
 FT FT STRAND 1122 1122
 FT FT TURN 1123 1123
 FT FT STRAND 1127 1131
 FT FT HELIX 1132 1134
 FT FT TURN 1135 1136
 FT FT STRAND 1137 1141
 FT FT TURN 1144 1145
 FT FT STRAND 1148 1152
 FT FT STRAND 1155 1158
 FT FT TURN 1159 1162
 FT FT STRAND 1163 1166
 FT FT TURN 1173 1178
 FT FT STRAND 1184 1185
 FT FT STRAND 1188 1188
 FT FT STRAND 1190 1190

Query Match 60.2%; Score 112; DB 1; Length 1314;
 Best Local Similarity 100.0%; Pred. No. 1.1e-07;
 Matches 2.; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 3
 GON1_RAT STANDARD; PRT: 92 AA.

Qy 1 FNNFTVSPKVSASHLE 21
 Db 946 FNNFTVSPKVSASHLE 966

RESULT 2
 PURA LEGPN STANDARD; PRT: 431 AA.

ID PURA LEGPN
 AC QBRN2;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Adenylosuccinate synthetase (EC 6.3.4.4) (IMP--aspartate ligase)
 (ADSS) (AMPases).
 GN PURA.
 OS Legionella pneumophila
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Legionellales;
 OC Legionellaceae; Legionella.
 OX NCBI_TaxID=446

RN SEQUENCE FROM N.A.
 RA Rankin S. Li Z., Isberg R.R.;
 RT "Macrophage induced genes of Legionella pneumophila: protection from reactive intermediates and solute imbalance during intracellular growth";
 RT Submitted (JAN-2002) to the EMBL/GenBank/DBJ databases.
 RL -|- FUNCTION: Plays an important role in the de novo pathway of purine nucleotide biosynthesis.
 CC -|- CATALYTIC ACTIVITY: GTP + IMP + L-aspartate = GDP + phosphate + adenylsuccinate.
 CC -|- COFACTOR: Binds 1 magnesium ion per subunit (By similarity).
 CC -|- PATHWAY: AMP biosynthesis; first committed step.
 CC -|- SUBUNIT: Homodimer (By similarity).
 CC -|- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
 CC -|- SIMILARITY: Belongs to the adenylosuccinate synthetase family.

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CC DR EMBL; AP409018; ANN00648; 1; ~.
 DR HAMAP; MF_00011; ~.
 DR InterPro; IPR01114; Asucc synthetase.
 DR Pfam; PF00709; Adenylylsuccinatase.
 DR Prodom; PDO01188; Asucc synthetase.
 DR TIGRFAMs; TIGR00084; Pufe; 1.
 DR PROSITE; PS00533; ADENYLYLSCUCCINATE; 1.
 DR KW Purine biosynthesis, Ligase; GTP-binding; Metal-binding; Magnesium.
 FT NP BIND 13 19 GTP (POTENTIAL).
 FT ACT SITE 141 141 BY SIMILARITY.
 FT ACT SITE 148 148 BY SIMILARITY.
 FT METAL 14 14 MAGNESIUM (BY SIMILARITY).
 FT METAL 41 41 MAGNESIUM (VIA CARBONYL OXYGEN) (BY SIMILARITY).

SQ SEQUENCE 431 AA; 47381 MW; D146C1PAED550774 CRC64;

Query Match 33.9%; Score 63; DB 1; Length 431;
 Best Local Similarity 36.4%; Pred. No. 0.25;
 Matches 12.; Conservative 7; Mismatches 14; Indels 0; Gaps 0;

Qy 1 FNNFTVSPKVSASHLE 33
 Db 168 YHNFVLIQFQPAVDLESLGESLQWAEELRP 200

DR Pfam: PF00446; GRH: 1.
 PRINTS: PRO1541; GONADOLIBRNI.
 DR PRO1541; GNKH: 1.
 KW Cleavage on pair of basic residues; Hormone; Amidation; Hypothalamus;
 KW Placenta; Signal; Pyrrolidone carboxylic acid.
 FT SIGNAL 1 23 PROGONADOLIBERIN I.
 FT CHAIN 24 92 GONADOLIBERIN I.
 FT PEPTIDE 24 33 PROLACTIN RELEASE-INHIBITING FACTOR I.
 FT PEPTIDE 37 92 APPEARS TO BE ESSENTIAL FOR BIOLOGICAL
 ACT_SITE 26 26 ACTIVITY.
 FT MOD_RES 24 24 PYRROLIDONE CARBOXYLIC ACID.
 FT MOD_RES 33 33 AMIDATION (G-34 PROVIDE AMIDE GROUP).
 SQ SEQUENCE 92 AA; 10500 MW; 494B5C64DA8A3EB3 CRC64;

Query Match 33.1%; Score 61.5%; DB 1; Length 92;
 Best Local Similarity 51.7%; Pred. No. 0.076; Indels 7; Gaps 2;
 Matches 15; Conservative 3; Mismatches 4; PRT; 1274 AA.

QY 12 VPKVSA-----HLEG-PSLHKSYGLRP 33
 :||: :|||: |||: |||: |||:
 Db 4 IPKLMIAAVVLLTCLEGGCSQWNSYGLRP 32

RESULT 4

BXPF_CLOBO	STANDARD;	PRT;	1274 AA.
ID BXPF_CLOBO			
AC P30596;			
DT 01-JUL-1993 (Rel. 26; Created)			
DT 01-JUL-1993 (Rel. 26; Last sequence update)			
DT 28-FEB-2003 (Rel. 41; Last annotation update)			
DE Botulinum neurotoxin type F precursor (EC 3.4.24.69) (BoNT/F)			
DS (Bontoxilysin F).			
GN Clostridium botulinum.			
OS Bacteria; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;			
OC Clostridium.			
OX NCBI_TaxID=1491;			
RN [1]			
RP SEQUENCE FROM N.A.			
RC STRAIN=ATCC 23387; PubMed=1398040;			
RD East A.K., Richardson P.T., Allaway D., Collins M.D.,			
RA Roberts T.A., Thompson D.B.;			
FT "Conserved structure of genes encoding components of Clostridium			
FT botulinum".			
RL PEMS Microbiol. Lett. 75:225-230(1992).			
RN [2]			
RP SEQUENCE OF 1-64 FROM N.A.			
RC STRAIN=Hobbs PT10;			
RA MEDLINE=94297488; PubMed=7764998;			
RA East A.K., Collins M.D.,			
RA "Conserved structure of genes encoding components of botulinum			
FT neurotoxin complex M and the sequence of the gene coding for the			
FT nontoxic component in nonproteolytic Clostridium botulinum type F.";			
RL Curr. Microbiol. 29:69-77(1994).			
RN [3]			
RP SEQUENCE OF 634-1002 FROM N.A.			
RA MEDLINE=94013372; PubMed=1408542;			
RA Campbell K., East A.K., Collins M.D.,			
RA "Gene probes for identification of the botulinum neurotoxin gene and			
FT specific probes for identification of neurotoxin types B, E, and F.";			
RL J. Clin. Microbiol. 31:2255-2262(1993).			
RN [4]			
RP IDENTIFICATION OF SUBSTRATE.			
RA MEDLINE=94230352; PubMed=175689;			
RA Yamasaki S., Baumheister A., Blasius J., Link E., Cornille F.,			
RA Roques B., Pyke P.M., Sudhof T.C., Jain R., Niemann H.,			
RA "Cleavage of members of the synaptobrevin/VAMP family by types D and			
FT F botulinum neurotoxins and tetanus toxin.";			
RL J. Biol. Chem. 269:12764-12772(1994).			
CC -!- FUNCTION: Botulinus toxin acts by inhibiting neurotransmitter			
CC RELEASE. IT BINDS TO PERIPHERAL NEURONAL SYNAPSES, IS INTERNALIZED			

NCBI_TaxID=1491;

RN [1] SEQUENCE FROM N.A.

RC STRAIN=13 / 30;

RC MEDLINE=94092745; PubMed=8268233;

RA Campbell K.; Collins M.D.; East A.K.;

RT "Nucleotide sequence of the gene coding for Clostridium botulinum (Clostridium argentinense) type G neurotoxin: genealogical comparison with other clostridial neurotoxins." [93].

RL Biochim. Biophys. Acta 1216:487-491 (1993).

CC -!- FUNCTION: BOTULINUM TOXIN ACTS BY INHIBITING NEUROTRANSMITTER RELEASE. IT BINDS TO PERIPHERAL NEURONAL SYNAPSIS. IS INTERNALIZED AND MOVES BY RETROGRADE TRANSPORT UP THE AXON INTO THE SPINAL CORD WHERE IT CAN MOVE BETWEEN POSTSYNAPTIC AND PRESYNAPTIC NEURONS. IT INHIBITS NEUROTRANSMITTER RELEASE BY ACTING AS A ZINC ENDOPeptidase.

CC -!- CATALYTIC ACTIVITY: Limited hydrolysis of Proteins of the neuracecytosis apparatus, synaptobrevins, SNAP25 or syntaxin. No detected action on small molecule substrates.

CC -!- COFACTOR: Binds 1 zinc ion per subunit (by similarity).

CC -!- SUBUNIT: Disulfide-linked heterodimer of a light chain (L) and a heavy chain (H). The light chain has the pharmacological activity, while the N- and C-terminal of the heavy chain mediate channel formation and toxin binding, respectively.

CC -!- SUBCELLULAR LOCATION: Secreted (By similarity).

CC -!- MISCELLANEOUS: There are seven antigenically distinct forms of botulinum neurotoxin: Types A, B, C1, D, E, F, and G.

CC -!- SIMILARITY: Belongs to peptidase family M27.

CC -----

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CC -----

CC EMBL; X74162; CBAA52275.1; -.

DR HSSP; P10845; 3BTA.

DR MBROPe; M27.002; -.

DR InterPro; IPR003985; ConA-like lec_g1.

DR InterPro; IPR003160; KunigZ_legume.

DR InterPro; IPR003025; Pept_M-Zn_BS.

DR InterPro; IPR003395; Peptidase_M27.

DR Pfam; PF01742; Peptidase_M27; 1.

DR PRINTS; PR00760; BONTOXIYLIN.

DR ProDom; PD001963; BONTOXIYLIN; 1.

DR PROSITE; PS00142; ZINC PROTEASE; 1.

DR Neurotoxin_Hydrolase_Metalloprotease_Zinc.

FT INIT_MET 0 0 BY SIMILARITY.

FT CHAIN 1 441 BOTULINUM NEUROTOXIN G, LIGHT-CHAIN.

FT METAL 229 229 ZINC (CATALYTIC) (BY SIMILARITY).

FT ACT_SITE 230 230 BY SIMILARITY.

FT METAL 233 233 ZINC (CATALYTIC) (BY SIMILARITY).

FT DISULFID_F 435 449 INTERCHAIN (PROBABLE).

SEQUENCE 1296 AA; 149013 MW; DCCE47E15F665C31 CRC64;

Query Match 32.8%; Score 61; DB 1; Length 1296;

Best Local Similarity 38.1%; Pred. No. 1.6;

Matches 9; Conservative 9; Mismatches 4; Indels 0; Gaps 0;

Qy 1 FNNFTVSFWLRYPKVASHLE 21

 | : | : | : | : | : | : ::

Db 929 FDNFNSINTWVTRPCKNNNDIQ 949

DT 28-PDB-2003 (Rel. 41, Last annotation update)

DE Botulinum neurotoxin type A precursor (EC 3.4.24.69). (BONT/A)

DE (Botoxylisin A) (BOTOK) [Contains: Botulinum neurotoxin A, light-chain, Botulinum neurotoxin A, heavy-chain].

DE BOTA OR BNA OR ATX.

GN Clostridium botulinum.

OS Bacteria; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;

OC Clostridium.

NCBI_TaxID=1491;

OX RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=NCTC 216;

RX MEDLINE=9023564; PubMed=2105020;

RA Thompson D.E.; Breton J.K.; Quillardet P.; Swinfield T.-J., Shone C.C.; Atkinson T.; Malling J.; Minton N.P.;

RT "The complete amino acid sequence of the Clostridium botulinum type A neurotoxin, deduced by nucleotide sequence analysis of the encoding gene." [94].

RT RL Biochem. 189:73-81 (1990).

RN RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=62A;

RX MEDLINE=90264400; PubMed=2160960;

RA Binz B.; Kuarzono H.; Wille M.; Freudenthal J.; Wernars K.; Niemann H.; RT "Organization and phylogenetic interrelations of genes encoding the complete sequence of botulinum neurotoxin type A and comparison with other clostridial neurotoxins." [95].

RT RL J. Biol. Chem. 265:9153-9158 (1990).

RN RN [3]

RP SEQUENCE OF 1-65 FROM N.A.

RC STRAIN=62A;

RX MEDLINE=9716817; PubMed=8863443;

RA East A.K.; Bhandari M.; Stacey J.M.; Campbell K.D.; Collins M.D.; RT "Organization and phylogenetic interrelations of genes encoding components of the botulinum toxin complex in protolysogenic Clostridium botulinum types A, B, and F: evidence of chimeric sequences in the gene encoding the nontoxic nonhemagglutinin component." [96].

RT RL Int. J. Syst. Bacteriol. 46:1105-1112 (1996).

RN RN [4]

RP SEQUENCE OF 1-34 FROM N.A.

RC STRAIN=Hall;

RX MEDLINE=89350959; PubMed=2669749;

RA Bettley M.J.; Somers E.; Dasgupta B.R.;

RT "Characterization of botulinum type A neurotoxin gene: delineation of the N-terminal encoding region." [97].

RT RL Biochem. Biophys. Res. Commun. 162:1388-1395 (1989).

RN RN [5]

RP SEQUENCE OF 1-18 FROM N.A.

RC STRAIN=Type A NIH;

RX MEDLINE=96036783; PubMed=8521962;

RA Fujita R.; Fujinaga Y.; Inoue K.; Nakajima H.; Kumon H.; Oguma K.; RT "Molecular characterization of two forms of nontoxic-nonhemagglutinin components of Clostridium botulinum type A progenitor toxins." [98].

RT RL FEBS Lett. 376:41-44 (1995).

RN RN [6]

RP SEQUENCE OF 1-16.

RX MEDLINE=4417501; PubMed=6370255;

RA Schmidt J.J.; Sathyamoorthy V.; Dasgupta B.R.;

RT "Partial amino acid sequence of the heavy and light chains of botulinum neurotoxin type A." [99].

RT RL Biochem. Biophys. Res. Commun. 119:900-904 (1984).

RN RN [7]

RP SEQUENCE OF 1-46.

RA Dasgupta B.R.; Foley J.; Niece R.;

RT "Partial sequence of the light chain of botulinum neurotoxin type A." [100].

RT RL Biochemistry 26:4162-4162 (1987).

RN RN [8]

RP SEQUENCE OF 1-5 AND 444-456.

RX MEDLINE=91120847; PubMed=2126206;

RA Dasgupta B.R.; Dekleva M.L.;

RT "Partial sequence of the light chain of botulinum neurotoxin type A: sequence of amino acids at the N-terminus and around the nicking site." [101].

RT RL Biochimie 72:661-664 (1990).

RN RN [9]

RESULT 6

BY1-CLOBO-CLOBO STANDARD; PRT; 1295 AA.

ID BY1-CLOBO-CLOBO

AC P018639; P01561; P148639;

DT 01-JUL-1989 (Rel. 1.11, Created)

DT 01-JUL-1993 (Rel. 1.26, Last sequence update)

SEQUENCE OF 448-464 AND 872-895.
 RX MEDLINE=89024462; PubMed=317218;
 RA Sathyamoorthy V.; Dasgupta B.R.; Foley J.; Niece R.L.;
 RT "Botulinum neurotoxin type A: cleavage of the heavy chain into two
 halves and their partial sequences." Arch. Biochem. Biophys. 266:142-151 (1988).
 [10]
 RN
 RP SEQUENCE OF 448-482.
 RX MEDLINE=85545016; PubMed=3896784;
 RA Shone C.C., Hambleton P., Meiling J.;
 RT and purification of Clostridium botulinum type A neurotoxin by trypsin
 and purification of two tryptic fragments. Proteolytic action near
 the COOH-terminal of the heavy subunit destroys toxin-binding
 activity";
 RL Eur. J. Biochem. 151:75-82 (1985).
 [11]
 RN RP IDENTIFICATION OF SUBSTRATE.
 RX MEDLINE=94063091; PubMed=8241676;
 RA Schiavo G., Santucci A., Dasgupta B.R., Mehta P.P., Jontes J.,
 Benfenati F., Wilson M.C., Montecucco C.;
 RT "Botulinum neurotoxins secreteys A and E cleave SNAP-25 at distinct
 COOH-terminal bonds." FEBS Lett. 335:9-13 (1993).
 RL
 RN RP IDENTIFICATION OF SUBSTRATE.
 RX MEDLINE=94124435; PubMed=8294407;
 RA Binz T., Blasi J., Yamasaki S., Baumeister A., Link E., Suedhof T.C.,
 Jahn R., Niemann H.;
 RA "Proteolysis of SNAP-25 by types E and A botulinum neurotoxins." J. Biol. Chem. 269:1617-1620 (1994).
 RN [13] RT
 RN [14] RT
 RX MEDLINE=21556941; PubMed=1700044;
 RA Rigoni M., Caccin P., Johnson E.A., Montecucco C., Rossetto O.;
 RT "Site-directed mutagenesis identifies active-site residues of the
 light chain of botulinum neurotoxin type A." J. Biol. Chem. 269:1617-1620 (1994).
 RN RP X-RAY CRYSTALLOGRAPHY (3.3 ANGSTROMS).
 RX MEDLINE=98455071; PubMed=9783750;
 RA Lacy D.B., Tepp W., Cohen A.C., Dasgupta B.R., Stevens R.C.;
 RT "Crystal structure of botulinum neurotoxin type A and implications
 for toxicity." Nat. Struct. Biol. 5:898-902 (1998).
 CC -|- FUNCTION: Inhibits acetylcholine release. The botulinum toxin
 binds with high affinity to peripheral neuronal presynaptic
 membrane, is then internalized by receptor-mediated endocytosis.
 CC The C-terminus of the heavy chain (H) is responsible for the
 CC adhesion of the toxin to the cell surface while the N-terminus
 CC mediates transport of the light chain from the endocytic vesicle
 CC to the cytosol. After translocation, the light chain (L)
 CC hydrolyses the 197-Gln-| Arg-198 bond in SNAP-25, thereby blocking
 CC neurotransmitter release. Inhibition of acetylcholine release
 CC results in flaccid paralysis, with frequent heart or respiratory
 CC failure.
 CC -|- CATALYTIC ACTIVITY: Limited hydrolysis of proteins of the
 CC neurosecretory apparatus, synaptobrevins, SNAP25 or syntaxin. No
 CC detected action on small molecule substrates.
 CC -|- COPACTOR: Binds 1 zinc ion per subunit.
 CC SUBUNIT: Disulfide-linked heterodimer of a light chain (L) and a
 CC heavy chain (H).
 CC -|- SUBCELLULAR LOCATION: Secreted.
 CC -|- PHARMACEUTICAL: Available under the name BOTOX (Allergan) for
 CC the treatment of strabismus and blepharospasm associated with
 CC dystonia and cervical dysinia. Also used for the treatment of
 CC hemifacial spasm and a number of other neurological disorders
 CC characterized by abnormal muscle contraction.
 CC -|- MISCELLANEOUS: There are seven antigenically distinct forms of
 CC botulinum neurotoxin: Types A, B, C1, D, E, F, and G.
 CC -|- SIMILARITY: Belongs to peptidase family M27.
 CC -|- DATABASE: NAME=BOTOX Product information Web site;
 CC WWW="http://www.botox.com/index.jsp?hp=productinfo".
 CC -|- DATABASE: NAME=Protein Spotlight;

NOTE=Issue 19 of February 2002;
 WWW="http://www.expasy.org/spc/light/articles/sptlt019.html".

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CC DR X52066 / CAA3629.1;
 CC DR M30196 / AAQ2322.1;
 CC DR EMBL / X92973 / CAA63551.1;
 CC DR EMBL / D67030 / BAA11051.1;
 CC DR EMBL / M27892 / ARA3269.1;
 CC DR PIR / A55294 / BTCLAB.
 CC DR PDB; 3BPA; 01-OCT-99.
 CC DR MEROPS; M27_0002/-.
 CC DR InterPro; IPR008385; ConA-like_lec_g1.
 CC DR InterPro; IPR002160; Runitz_legume.
 CC DR InterPro; IPR006026; PepT_M_Zn_BS.
 CC DR InterPro; IPR00395; PepIDase_M27.
 CC DR Pfam; PF01742; Peptidase_M27_1.
 CC DR PRINTS; PR00760; BONTOKYNSIN.
 CC DR PRODOM; PD001963; BONTOKYNSIN.
 CC DR PROSITE; PS00112; ZINC_Protease_1.
 CC KW Neurotoxin; Transmembrane_Hydrolase; Metalloprotease; Zinc; Pharmaceutical; 3D-structure.

CC FT INIT_MET 0 0
 CC FT CHAIN 1 447
 CC FT CHAIN 448 1295
 CC FT MEVAL 222 222
 CC FT ACT_SITE 223 223
 CC FT METAL 226 226
 CC FT METAL 261 261
 CC FT DISULFID 429 453
 CC FT DISULFID 1234 1279
 CC FT TRANSMEM 626 646
 CC FT TRANSMEM 655 675
 CC FT VARIANT 26 26
 CC FT MUTAGEN 261 261
 CC FT MUTAGEN 265 265
 CC FT MUTAGEN 365 365
 CC FT CONFLICT 1 1
 CC FT CONFLICT 479 479
 CC FT CONFLICT 875 875
 CC FT CONFLICT 891 891
 CC SQ SEQUENCE 1295 AA; 149322 MW; 958342F54862579 CRCS4;
 CC
 CC Query Match 31.7%; Score 59; DB 1; Length 1295;
 CC Best Local Similarity 57.1%; Pred. No. 3;
 CC Matches 8; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

RESULT 7
 BXE_CLOBO STANDARD; PRT; 1250 AA.
 ID BXE_CLOBO
 AC Q00196;
 DT 01-JUL-1993 (Rel. 26, Created)
 DT 01-JUL-1993 (Rel. 26, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DB Botulinum neurotoxin type B precursor (EC 3.4.24.69) (BONT/B)
 DE (Botoxilysin B).
 OS Clostridium botulinum.
 OC Bacillus; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;
 OC Clostridium.
 NCBI_TaxID=1491;

QY 1 FNNTVSEMLRK 14
 : ||: ||: ||: ||:
 Db 937 YENFTSFWRIPK 950

- [1] SEQUENCE FROM N. A.
 RP STRAIN=Bela;
 RX MEDLINE=92181428; PubMed=1543481;
 RA Poulet S., Hauss D., Quanz M., Niemann H., Popoff M.R.; Clostridium
 "Sequencing of the botulinum neurotoxin E derived from Clostridium
 botulinum type E (strain Belga) and Clostridium butyricum (strains
 ATCC 43181 and ATCC 43755)." ;
 RL Biochem. Biophys. Res. Commun. 183:107-113 (1992).
 RN [2] SEQUENCE FROM N. A.
 RP MEDLINE=92174922; PubMed=1541280;
 RA Whelan S.M., Elmore M.J., Bodsworth N.J., Atkinson T., Minton N.P.;
 "The complete amino acid sequence of the Clostridium botulinum type-E
 neurotoxin, derived by nucleotide-sequence analysis of the encoding
 gene." ;
 RL Eur. J. Biochem. 204:657-667(1992).
 RN [3] SEQUENCE OF 1-251 FROM N. A.
 RP MEDLINE=90266400; PubMed=160960;
 RA Binz T., Kurazono H., Wille M., Frevert J., Wernars K., Niemann H.;
 "The complete sequence of botulinum neurotoxin type A and comparison
 with other clostridial neurotoxins." ;
 RL J. Biol. Chem. 265:9153-9158(1990).
 RN [4] SEQUENCE OF 1-13.
 RX MEDLINE=85191963; PubMed=3888113;
 RA Schmidt J.-U., Sathyamoorthy V., Dasgupta B.R.; "Botulinum neurotoxins types B and
 partial amino acid sequences of botulinum neurotoxins types B and
 E;" ;
 RT Arch. Biochem. Biophys. 238:544-548 (1985).
 RN [5] SEQUENCE OF 419-426.
 RP MEDLINE=90344918; PubMed=2116911;
 RA Gómez J.A., Dasgupta B.R.; "Botulinum neurotoxin type E fragment with endoprotease Lys-C
 reveals the site trypsin nicks and homology with tetanus
 neurotoxin." ;
 RL Biochimie 72:213-217(1990).
 RN [6] IDENTIFICATION OF SUBSTRATE.
 RX MEDLINE=94063091; PubMed=8243676;
 RA Schiavo G., Santucci A., Dasgupta B.R., Mehta P.P., Jontes J.,
 Bentivoglio F., Wilson M.C., Montecucco C.; "Botulinum neurotoxins serotypes A and E cleave SNAP-25 at distinct
 COOH-terminal peptide bonds." ;
 RL FEBS Lett. 335:99-103 (1993).
 RN [7] IDENTIFICATION OF SUBSTRATE.
 RP MEDLINE=94124495; PubMed=8224407;
 RX Binz T., Blasi J., Yamasaki S., Baumeister A., Link E., Suedhof T.C.,
 RA Jahn R., Niemann H.; "Proteolysis of SNAP-25 by types B and A botulinum neurotoxins." ;
 RT "J. Biol. Chem. 269:1617-1620(1994)."
 RL "- FUNCTION: BOTULINUM TOXIN ACTS BY INHIBITING NEUROTRANSMITTER
 RELEASE IT BINDS TO PERIPHERAL NEURONAL SYNAPSES, IS INTERNALIZED
 AND MOVES BY RETROGRADE TRANSPORT UP THE AXON INTO THE SPINAL CORD
 WHERE IT CAN MOVE BETWEEN POSTSYNAPTIC AND PRESYNAPTIC NEURONS. IT
 INHIBITS NEUROTRANSMITTER RELEASE BY ACTING AS A ZINC
 ENDOPPIETIDE THAT CATALYZES THE HYDROLYSIS OF THE 180-ARG-|-ILE-
 181 BOND IN SNAP-25.
 CC -|- CATALYTIC ACTIVITY: Limited hydrolysis of proteins of the
 CC neuroexocytosis apparatus, synaptobrevin, SNAP25 or syntaxin. No
 CC detected action on small molecule substrates.
 CC -|- COFACTOR: Binds 1 zinc ion per subunit (By similarity).
 CC -|- SUBUNIT: Disulfide-linked heterodimer of a light chain (L) and a
 CC heavy chain (H). The light chain has the pharmacological activity,
 CC while the N- and C-terminal of the heavy chain mediate channel
 CC formation and toxin binding, respectively.
 CC -|- SUBCELLULAR LOCATION: Secreted.
 CC -|- MISCELLANEOUS: There are seven antigenically distinct forms of
 CC botulinum neurotoxin: Types A, B, C1, D, E, F, and G.
 CC -|- SIMILARITY: Belongs to peptidase family M27.
- [1] This SWISS-PROT entry is copyright. It is produced through a collaboration
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 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC ---
 DR EMBL; X62089; CAA43998; 1; -.
 DR EML; X62683; CAA4556; 1; -.
 DR PIR; S08575; S08575.
 DR P21178; S21178.
 DR HSSP; P1045; 3BPA.
 DR MEOPS; M27_002; -.
 DR InterPro; IPR008385; ConA-like lec_g1.
 DR InterPro; IPR021260; Kunz_F_leucine.
 DR InterPro; IPR006025; Pept_M_Zn_BS.
 DR InterPro; IPR000395; Peptidase_M27.
 DR Pfam; PRO0760; Peptidase_M27_1.
 DR PRNTS; PRO00963; BONTXXLISIN.
 DR ProDom; PD00963; Bontoxilysin_1.
 DR PROSITE; PS00142; ZINC PROTEASE_1.
 DR Neurotoxin; Transmembrane_Hydrolase; Metalloprotease; Zinc.
 KW KW Neurotoxin; Transmembrane_Hydrolase; Metalloprotease; Zinc.
 DR INIT MET 0 0
 FT CHAIN 1 421 BOTULINUM NEUROTOXIN_E, LIGHT-CHAIN.
 FT CHAIN 422 1250 BOTULINUM NEUROTOXIN_E, HEAVY-CHAIN.
 FT METAL 211 211 ZINC (CATALYTIC) (BY SIMILARITY).
 FT ACT SITE 212 212 BY SIMILARITY.
 FT METYL 215 215 ZINC (CATALYTIC) (BY SIMILARITY).
 FT DISULFID 411 425 INTERCHAIN (PROBABLE).
 FT CONFLICT 176 176 R -> G (IN REF. 2).
 FT CONFLICT 197 197 S -> S (IN REF. 2 AND 3).
 FT CONFLICT 339 339 R -> A (IN REF. 2).
 FT CONFLICT 772 772 I -> L (IN REF. 2).
 FT CONFLICT 962 963 FE -> LQ (IN REF. 2).
 FT CONFLICT 966 966 R -> A (IN REF. 2).
 FT CONFLICT 1194 1194 N -> NN (IN REF. 2).
 SQ SEQUENCE 1250 AA; 143712 MW; D9FC26DDA041EB4 CRC64;
- Query Match 31.5%; Score 58.5%; Length 1250;
 Best Local Similarity 23.6%; Pred. No. 3.4%;
 Matches 13; Conservative 9; Mismatches 4; Indels 29; Gaps 2;
- Qy 1 FNNFTVFSWLRVP-----KVSASHLGPSLHNSY 29
 Db 911 YKNNFSFVNTRIPYDNKIVNVNEYLTINCRDNNSGMKVSNHNE--IIWF 962
- RESULT 8
 BXB_CLOBO
 ID BXB_CLOBO STANDARD; PRT; 1290 AA.
 AC P10844; P10843;
 DT 01-JUL-1999 (Ref. 11, Created)
 DT 01-OCT-2003 (Ref. 26, Last sequence update)
 DT 10-OCT-2003 (Ref. 42, Last annotation update)
 DE Botulinum neurotoxin type B precursor (EC 3.4.24.69) (BoNT/B).
 DE (Bontoxilysin B).
 GN
 OS Clostridium botulinum. Clostridia; Clostridiales; Clostridiaceae;
 OC Bacteria; Firmicutes;
 OC Clostridium.
 RN [1]
 RN NCBI_TaxId=1491;
 RN
 RP SEQUENCE FROM N.A.
 RX MEDLINE=92384550; PubMed=1514783;
 RA Whelan S.M., Elmore M.J., Bodsworth N.J., Brehm J.K., Atkinson T.,
 RA Minton N.P.;
 RT "Molecular cloning of the Clostridium botulinum structural gene
 encoding the type B neurotoxin and determination of its entire
 RT nucleotide sequence." ;
 RL Environ. Microbiol. 58:2345-2354 (1992).
 RN

SEQUENCE OF 35-245 FROM N.A.
 STRAIN=NCTC 7273;
 RA Szabo E.A., Pemberton J.M., Desmarchelier P.N.; databases.
 RL Submitted (AFR-1992) to the EMBL/Genbank/DBJ databases.
 RN [3]
 SEQUENCE OF 633-993 FROM N.A.
 RP STRAIN=CTC 7273;
 RX MEDLINE=94013372; PubMed=8408542;
 RA Campbell K., East A.K., Collins M.D.;
 RT "Gene probes for identification of the botulinum neurotoxin gene and
 specific identification of neurotoxin types B, E, and F.";
 RL J. Clin. Microbiol. 31:2255-2262(1993).
 [4]
 SEQUENCE OF 1-44 AND 441-466.
 RP STRAIN=657;
 RX MEDLINE=89000987; PubMed=3139097;
 RA Dasgupta B.R., Datta A.;
 RT "Botulinum neurotoxin type B (strain 657): partial sequence and
 similarity with tetanus toxin";
 RL Biochimie 70:811-817(1988).
 [5]
 SEQUENCE OF 1-16 AND 441-458.
 RP STRAIN=ORRA;
 RX MEDLINE=85197963; PubMed=3888113;
 RA Schmidt J.J., Sathyamoorthy V., Dasgupta B.R.;
 RT "Partial amino acid sequences of botulinum neurotoxins types B and
 E.";
 RL Arch. Biochem. Biophys. 238:544-548(1985).
 [6]
 RN IDENTIFICATION AS ZINC-PROTEASE.
 RX MEDLINE=93054694; PubMed=1422690;
 RA Schiavo G., Rossetto O., Santucci A., Dasgupta B.R., Montecucco C.;
 RT "Botulinum neurotoxins are zinc proteins.";
 RL J. Biol. Chem. 271:23479-23483(1998).
 [7]
 RP IDENTIFICATION OF SUBSTRATE.
 RX MEDLINE=93063293; PubMed=13131807;
 RA Schiavo G., Benfenati F., Poulain B., Rossetto O., de Laureto P.P.,
 RT "Tetanus and botulinum-B neurotoxins block neurotransmitter release
 by proteolytic cleavage of synaptobrevin.";
 RL Nature 359:832-835(1992).
 CC FUNCTION: BOTULINUM TOXIN ACTS BY INHIBITING NEUROTRANSMITTER
 RELEASE. IT BINDS TO PERIPHERAL NEURONAL SYNAPSES. IS INTERNALIZED
 AND MOVES BY RETROGRADE TRANSPORT UP THE AXON INTO THE SPINAL CORD
 WHERE IT CAN MOVE BETWEEN POSTSYNAPTIC AND PRESYNAPTIC NEURONS. IT
 INHIBITS NEUROTRANSMITTER RELEASE BY ACTING AS A ZINC
 ENDOPePTIDase THAT CLEAVES THE 76-GUN-1-PHE-77 BOND OF
 SYNAPTOBREVIN-2.
 CC CATALYTIC ACTIVITY: Limited hydrolysis of proteins of the
 neuroexocytosis apparatus, synaptobrevins, SNAP25 or syntaxin. No
 CC DEFECTED action on small molecule substrates.
 CC SUBUNIT: Disulfide-linked heterodimer of a light chain (L) and a
 heavy chain (H). The light chain has the pharmacological activity,
 formation and toxin binding, respectively.
 CC SUBCELLULAR LOCATION: Secreted.
 CC - MISCELLANEOUS: There are seven antigenically distinct forms of
 botulinum neurotoxin: Types A, B, C1, D, E, F, and G.
 CC - SIMILARITY: Belongs to Peptidase family M27.

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 modified and this statement is not removed. Usage by and for commercial
 entities requires license agreement (see <http://www.isb-sib.ch/announce/>
 or send an email to licenses@isb-sib.ch).

CC DR EMBL; M81186; AA123211.1;
 CC DR EMBL; Z11934; CAA7991.1;
 CC DR EMBL; X70817; CAA50148.1;
 CC DR EMBL; 211934; CAA7991.1;
 CC DR EMBL; X70817; CAA50148.1;

CC -1- SUBCELLULAR LOCATION: Secreted.
 CC -1- SIMILARITY: Belongs to the GrnRH family.

DR PIR; A48940; A48940.
 DR IBEW; 01-NOV-00.
 DR PDB; 1F31; 01-NOV-00.
 DR PDB; 1FB2; 16-AUG-00.
 DR PDB; 1FB3; 16-AUG-00.
 DR PDB; 1FOH; 06-DEC-00.
 DR PDB; 1GFB; 13-NOV-02.
 DR PDB; 1G9C; 13-NOV-02.
 DR PDB; 1IGD; 13-NOV-02.
 DR PDB; 1IIB; 21-NOV-01.
 DR InterPro; IP008985; Cona like lec_g1.
 DR InterPro; IP002160; KunifIZ legume.
 DR InterPro; IP006025; KunifIZ legume.
 DR InterPro; IP000355; Peptidase_M27.
 DR Pfam; PF01742; Peptidase_M27; 1.
 DR PRINTS; PRO0760; BONTOKIYTSN.
 DR ProDom; PD001963; Botocoxilyein; 1.
 DR PROSITE; PS00142; ZINC PROTEASE; 1.
 DR Neurotoxin; Transmembrane; Hydrolase; Metalloprotease; Zinc; KW 3D-structure.
 KW INIT MET 0 0
 FT CHAIN B, LIGHT-CHAIN 0 0
 FT CHAIN A, HEAVY-CHAIN 0 0
 FT METAL 440 440
 FT ACT SITE 1290 1290
 FT ACT SITE 229 229
 FT ACT SITE 230 230
 FT ACT SITE 233 233
 FT ACT SITE 233 233
 FT DISULFID 436 436
 FT CONFLICT 45 45
 FT CONFLICT 29 29
 FT CONFLICT 217 217
 FT CONFLICT 224 224
 FT CONFLICT 463 463
 FT CONFLICT 5 5
 SEQUENCE 1290 AA; 150670 MW; D21746E2CO4DF43 CRC64;
 Query Match 31 2%; Score 58; DB 1; Length 1290;
 Best Local Similarity 64 3%; Pred. No. 4.1;
 Matches 9; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Qy 1 FNNFTYSFWLVRPK 14
 Db 922 FLDFYSFWLVRPK 935

RESULT 9
 ID GON1_MOUSE STANDARD; PRT;
 AC P13562;
 DT 01-JAN-1990 (Rel. 13, Created)
 DT 01-JAN-1990 (Rel. 13, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Progonadotropin I precursor (Contains: Gonadotropin I (LH-RH I))
 DE Luteinizing hormone releasing hormone I (Gonadotropin-releasing factor
 DE hormone I) (GrnRH I) (Luteinizing hormone I); Prolactin release-inhibiting factor
 DE 1);
 GN GRNHL OR GRNHL
 OS Mus musculus (Mouse)
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Buteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Murinae; Mus.
 OC NCBI_TAXID:10090;
 RN [L]
 RP SEQUENCE FROM N.A.
 RX MEDLINE:87059928; PubMed=3024317;
 RA Mason A.J., Haylick J.S., Zoeller R.T., Young W.S. III,
 RA Phillips H.S., Nikolicic K., Seeburg P.H.;
 RT "A deletion truncating the gonadotropin-releasing hormone gene is
 responsible for hypogonadism in the hpg mouse.";
 RT Science 234:1366-1371(1986).
 CC -1- FUNCTION: Stimulates the secretion of gonadotropins; it stimulates
 CC the secretion of both luteinizing and follicle-stimulating
 CC hormones.
 CC -1- SUBCELLULAR LOCATION: Belongs to the GrnRH family.
 CC -1- SIMILARITY: Belongs to the GrnRH family.

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EMBL; M14872; AAA37717.1; -.

PIR; A47578; RIMSG.

MGD; MGI:5789;

InterPro; IPR004012; GnRH.

DR InterPro; IPR004079; Gonadotropin.

DR PRINTS; PRO1541; GONADOLIBERNI.

DR PROSITE; PS00474; GnRH_1.

KW Cleavage on pair of basic residues; Hormone; Amidation; Hypothalamus; Placenta; Signal; Pyrrolidone carboxylic acid.

FT SIGNAL 1 21 PROGONADOLIBERIN_I.

FT PEPTIDE 22 90 GONADOLIBERIN_I.

FT PEPTIDE 35 90 PROLACTIN RELEASE-INHIBITING FACTOR_I.

FT ACT_SITE 24 24 APPEARS TO BE ESSENTIAL FOR BIOLOGICAL ACTIVITY.

FT MOD_RES 22 22 PYRROLIDONE CARBOXYLIC ACID.

FT MOD_RES 31 31 AMIDATION (G-32 PROVIDE AMIDE GROUP) .

SQ SEQUENCE 90 AA; 10337 MW; 1C076FA4826ED9 CRC64;

Query Match Score 57.5; DB 1; Length 90; Best Local Similarity 80.0%; Pred. No. 0.27; Matches 12; Conservative 0; Mismatches 2; Indels 1; Gaps 1; Qy 20 LEG-PSLHWSYGLRP 33 Db 16 LEGCSSOHWSYGLRP 30

RESULT 10 VP2_AHSV6 STANDARD: PRT; 1051 AA.

AC 071024; 15-DEC-1998 (Rel. 37, Created)

DT 15-DEC-1998 (Rel. 42, Last sequence update)

DT 10-OCT-2003 (Rel. 42, Last annotation update)

DB Outer capsid protein VP2.

GN S2 OR L2.

OS African horse sickness virus 6 (AHSV-6) (African horse sickness virus (serotype 6)).

OS Viruses; dsRNA viruses; Reoviridae; Orbivirus.

OX NCBI_TAXID=86060;

RN SRQSEQUENCE FROM N.A.; PubMed=9617769;

RX Williams C.F., Itoue T., Lucius A.-M., Zanotto P., Roy P.,

RT "The complete sequence of four major structural proteins of African horse sickness virus serotype 6: evolutionary relationships within and between the orbiviruses.";

RT Virus Res. 53:53-73 (1998).

CC -!- FUNCTION: THE VP2 PROTEIN IS ONE OF THE TWO PROTEINS (WITH VP5) WHICH CONSTITUTE THE VIRUS PARTICLE OUTER CAPSID. IT IS THE MAJOR TARGET OF THE HOST IMMUNOCENIC RESPONSE.

CC -!- SIMILARITY: Belongs to the reoviruses VP2 protein family.

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CC AF021235; AAC40994.1; -.

DR InterPro; IPR001742; Orbi_VP2.

DR Pfam; PF00398; Orbi_VP2; I.

KW Coat Protein.

SQ SEQUENCE 1051 AA; 122226 MW; 2B04DB9E389F4B5F CRC44;

Query Match Score 57; DB 1; Length 1051; Best Local Similarity 47.6%; Pred. No. 4.6%; Matches 10; Conservative 4; Mismatches 7; Indels 0; Gaps 0;

Qy 1 FNNFTIVSFWLKPVKVSASHLE 21

Db 636 FSKRFVSYWYRVEKITTKHL 656

RESULT 11 GON1_RANCA STANDARD: PRT; 90 AA.

ID GON1_RANCA

AC Q9Y63;

DT 10-OCT-2003 (Rel. 42, Created)

DT 10-OCT-2003 (Rel. 42, Last sequence update)

DE Progonadotropin I precursor (Contains: Gonadotropin I (LHRH_I); Luteinizing hormone releasing hormone I (LHRH_I); Gonadotropin releasing hormone I (GnRH_I); GnRH-associated peptide I (GAP1)).

DE (Luteinizing hormone I (LHRH_I)) (Luteinizing hormone I (GnRH_I))

GN GNRH1 OR GNRH.

OS Ranctesbeiana (Bull frog).

OC Eukaryota; Metazoa; Chordata; Vertebrates; Euteleostomi; Amphibia; Batrachia; Anura; Neobatrachia; Ranidae; Rana.

OC NCBI_TAXID=8400;

RN [1] SOURCE FROM N.A., TISSUE SPECIFICITY, AND DEVELOPMENTAL STAGE.

RN TISSUE:Forebrain; RC MEDLINE:21102951; PubMed=11170016;

RX Wang L., Yoo M.S., Kang H.M., Im W.B., Choi H.S., Bogerd J., RA Kwon H.B.; RT "Cloning and characterization of cDNAs encoding the GnRH1 and GnRH2 precursors from bullfrog (Rana catesbeiana)."; RL J. Exp. Zool. 288:190-201(2001).

CC -!- FUNCTION: Stimulates the secretion of gonadotropins (By similarity).

CC -!- SUBCELLULAR LOCATION: Secreted.

CC -!- TISSUE SPECIFICITY: Forebrain.

CC -!- DEVELOPMENTAL STAGE: Expressed at significantly higher levels during post-embryonic development.

CC -!- SIMILARITY: Belongs to the GnRH family.

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CC CC -!- SUBCELLULAR LOCATION: Secreted.

CC -!- TISSUE SPECIFICITY: Forebrain.

CC -!- DEVELOPMENTAL STAGE: Expressed at significantly higher levels during post-embryonic development.

CC -!- SIMILARITY: Belongs to the GnRH family.

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CC DR EMBL; AF18874; AAU05912.1; GO: GO:0005576; C:extracellular; NAS.

DR GO: GO:0005153; P:G-protein coupled receptor activity; NAS.

DR GO: GO:0009755; P:hormone mediated signaling; NAS.

DR GO: GO:0000003; P:reproduction; NAS.

DR InterPro; IPR002012; GRH

DR InterPro; IPR004079; Gonadotropin.

DR Pfam; PF00446; GRH; I.

DR PRINTS; PRO1541; GONADOLIBERNI.

DR PROSITE; PS00473; GNRH_1.

DR Cleavage on pair of basic residues; Hormone; Amidation; Signal;

KW Pyrrolidone carboxylic acid. POTENTIAL.

FT SIGNAL 1 24 PROGONADOLIBERIN_I.

FT CHAIN 25 90 GONADOLIBERIN_I.

FT PEPTIDE 25 34 GNRH-ASSOCIATED PEPTIDE_I (BY SIMILARITY).

FT PEPTIDE 38 86 PYRROLIDONE CARBOXYLIC ACID (BY SIMILARITY).

FT MOD_RES 25 25 PT

- I- botulinum types A, B, and F: evidence of chimeric sequences in the gene encoding the nontoxic nonhemagglutinin component. ;
- I- Int. J. Syst. Bacteriol. 46:1105-1112(1996).
- I- FUNCTION: Inhibits acetylcholine release. The botulinum toxin binds with high affinity to peripheral neuronal presynaptic membrane, is internalized by receptor-mediated endocytosis. The C-terminus of the heavy chain (H) is responsible for the adherence of the toxin to the cell surface while the N-terminus mediates transport of the light chain from the endocytic vesicle to the cytosol. After translocation, the light chain (L) hydrolyzes the 19-Gln-1 Arg-198 bond in SNAP-25, thereby blocking neurotransmitter release. Inhibition of acetylcholine release results in flaccid paralysis, with frequent heart or respiratory failure (By similarity).
- I- CATALYTIC ACTIVITY: Limited hydrolysis of proteins of the neuroexocytosis apparatus, synaptobrevin, SNAP25 or syntaxin, detected action on small molecule substrates.
- I- SUBUNIT: Disulfide linked heterodimer of a light chain (L) and heavy chain (H) (By similarity).
- I- SUBCELLULAR LOCATION: Secreted.
- I- MISCELLANEOUS: There are seven antigenically distinct forms of botulinum neurotoxin: Types A, B, C1, D, E, F, and G.
- I- SIMILARITY: Belongs to Peptidase family M27.

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28-FEB-2003 (Rel. 41, Last annotation update)
 Progonadotropin I precursor [contains: Gonadal (luteinizing hormone I) (Gona hormone I); GnRH I] (Luliberin I) (Gonadorelin peptide I].

[1] SEQUENCE FROM N.A.
 Homo sapiens (Human).
 Metacercaria; Chordata; Craniata; Vertebrates;
 Mammalia; Eutheria; Primates; Catarrhini; Hominoidea;
 GNRH1 OR GnRH OR LHRH.

[2] SEQUENCE FROM N.A.
 MEDLINE=8916668; PubMed=671939;
 Hayflick J.S., Adelman J.P., Seeburg P.H.;
 "The complete nucleotide sequence of the human
 hormone gene.";
Nucleic Acids Res. 17:6403-6403 (1989).

[3] SEQUENCE FROM N.A., AND VARIANT SER-1-6.
 MEDLINE=85012739; PubMed=690951;
 Seeburg P.H., Adelman J.P.;
 "Characterization of cDNA for precursor of human
 releasing hormone.";

EMBL; X01059; CAA25526.1; -;
 DR EMBL; M12578; AA335916.1; -;
 DR EMBL; X15215; CA33285.1; -;
 DR PIR; S03308; RHTDG.
 DR Genew; HGNC; 4419; GNRHL.
 MTM; 15276; -;
 DR GO; GO:0005645; C:soluble fraction; TAS.
 DR GO; GO:0005133; Flutelinating hormone-releasing factor activity; TAS.
 DR GO; GO:0007257; P:cell-cell signaling; TAS.
 DR GO; GO:0008285; P:developmental; TAS.
 DR GO; GO:0007165; P:negative regulation of cell proliferation; TAS.
 DR InterPro; IPR00212; GrbH.
 DR InterPro; IPR04079; Gonadoliberin.
 DR PRINTS; PRO1541; GONADOLIBRNL.
 DR PROSITE; PS00473; GNRH_1.
 DR Cleavage on pair of basic residues; Hormone; Amidation; Hypothalamus;
 KW Placenta; Pharmaceutical; Signal; Polymorphism; Transduction; TAS.
 KW Pyrrolidone carboxylic acid.
 PT SIGNAL 1 23
 PT CHAIN 24 92
 PT PEPTIDE 24 33
 PT PEPTIDE 37 92
 FT ACT_SITE 26 26
 PT MOD_RES 24 24
 PT MOD_RES 33 33
 PT VARIANT 16 16
 PT AMIDATION (G-34 PROVIDE AMIDE GROUP).
 W -> S (in dBDNP:185).
 /FTid=VAR_013943.
 SQ SEQUENCE 92 AA; 10380 MW; 30AT221E076FA79 CRC64;
 Query Match 29 3%; Score 54.5; DB 1; Length 92;
 Best Local Similarity 73.3%; Pred. No. 0.73%; Indels 1; Gaps 1;
 Matches 11; Conservative 1; Mismatches 2; Indels 1; Gaps 1;

QY 20 LEG-PSIHWSYGLRP 33
 Db 18 VEGCQQHQWYGLRP 32

RESULT 15
 BXCL_CLOBO
 ID BXCL_CLOBO
 AC P18640;
 DT 01-NOV-1990 (Rel. 16, Created)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Botulinum neurotoxin type C1 precursor (EC 3.4.24.69) (BONT/C1)
 DE (Bontoxilysin C1)
 OS Clostridium botulinum.
 OC Closridium; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;
 OX NCBI_TAXID:1491
 RN [1] SEQUENCE FROM N.A.
 RX MEDLINE=90370487; PubMed=2204031;
 RA Hauser D., Eklund M.W., Kurazona H., Binz T., Niemann H., Gill D.M., Boquet P., Popoff M.R.;
 RT "Nucleotide sequence of Clostridium botulinum C1 neurotoxin.";
 RL Nucleic Acids Res. 18:4924-4924(1990).
 RN [2] SEQUENCE FROM N.A.
 RP STRAIN-TYPE C Stockholm / C-ST;
 RX MEDLINE=91024998; PubMed=222445;
 RA Kimura K., Fujii N., Tsuzuki K., Murakami T., Inoh T., Yokosawa N., Takeshi K., Syuto B., Oguma K.;
 RT "The complete nucleotide sequence of the gene coding for botulinum type C1 toxin in the C-ST phage genome.";
 RL Biochem. Biophys. Res. Commun. 171:1304-1311(1990).
 RN [3] SEQUENCE OF 2-25.
 RP STRAIN-Type C Stockholm / C-ST;

RX MEDLINE=88153072; PubMed=2450068;
 RA Tsuzuki K., Yokosawa N., Syuto B., Ohishi I., Fujii N., Niemann H., Kimura K., Oguma K.;
 RA "Establishment of a monoclonal antibody recognizing an antigenic site common to Clostridium botulinum type B, C1, D, and E toxins and tetanus toxin.";
 RT Infec. Immun. 56:1898-902(1988).
 RL [4]

RN FUNCTION OF SUBSTRATE.
 RP IDENTIFICATION: PubMed=701002;
 RX IDENTIFICATION: PubMed=94038966;
 RA Blasi J., Chapman E.R., Yamasaki S., Binz T., Niemann H., Jahn R., Boulinum neurotoxin C1 blocks neurotransmitter release by means of cleaving HPC-1/syntaxin.;
 RT EMBO J. 12:4821-4828(1993).

CC -|- FUNCTION: BOTULINUM TOXIN ACTS BY INHIBITING NEUROTRANSMITTER RELEASE. IT BINDS TO PERIPHERAL NEURONAL SYNAPSES, IS INTERNALIZED AND MOVES BY RETROGRADE TRANSPORT UP THE AXON INTO THE SPINAL CORD WHERE IT CAN MOVE BETWEEN POSTSYNAPTIC AND PRESYNAPTIC NEURONS. IT INHIBITS NEUROTRANSMITTER RELEASE BY ACTING AS A ZINC ENDOPeptidase THAT CLEAVES SYNTAXIN.

CC -|- CATALYTIC ACTIVITY: Limited hydrolysis of proteins of the neuroexocytosis apparatus, synaptobrevins, SNAP25 or syntaxin. No detected action on small molecule substrates.

CC -|- COFACTOR: Binds 1 zinc ion per subunit (By similarity).
 CC -|- SUBUNIT: Disulfide-linked heterodimer of a light chain (L) and a heavy chain (H). The light chain has the pharmacological activity, while the N- and C-terminal of the heavy chain mediate channel formation and toxin binding, respectively.

CC -|- SUBCELLULAR LOCATION: Secreted.

CC -|- MISCELLANEOUS: There are seven antigenically distinct forms of botulinum neurotoxins: Types A, B, C1, D, E, F, and G.

CC -|- MISCELLANEOUS: BOTULINUM TYPE C1 NEUROTOXIN IS SYNTHESIZED BY C STRAIN OF CLOSTRIDIUM BOTULINUM WHICH CARRY THE APPROPRIATE BACTERIOPHAGE.

CC -|- SIMILARITY: Belongs to peptidase family M27.

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CC EMBL; X66433; CAA41060.1; -;
 DR EMBL; X2293; CAA52313.1; -;
 DR EMBL; X53751; CAA37780.1; -;
 DR EMBL; D90210; BAA11235.1; -;
 DR EMBL; X62389; CAA44263.1; -;
 DR RSSP; P10845; 3BTA.
 DR MEROPS; M27.002; -;
 DR InterPro; IPR008985; ConA_like_lec_g1.
 DR InterPro; IPR00160; Kunitz_legume.
 DR InterPro; IPR000395; Pept_M_Zn_BS.
 DR InterPro; IPR000395; Peptidase_M27.
 DR PRINTS; PR00760; Bontoxilysin.
 DR ProDom; PD001963; Bontoxilysin.
 DR PROSITE; PS00142; ZINC_ProteaseB_1.
 DR INIT MET 0 0
 FT CHAIN 1 448
 FT CHAIN 0 1290
 FT METAL 228 228
 FT ACT SITE 229 229
 FT METAL 232 232
 FT DISULFID 436 452
 FT CONFLICT 84 84
 SQ SEQUENCE 1290 AA; 148734 MW; 71FBEB379F97129EB CRC64;

Query Match 29 3%; Score 54.5; DB 1; Length 1290;
 Best Local Similarity 30.8%; Pred. No. 13; Mismatches 10; Indels 9; Gaps 2;

Qy 1 FNNFTVFWLRVPKVASHLEGPSL-----HWSYGL 31
: : | : | : | : | : | : | : | : | : | : | : | : | :
Db 934 YESPSTFWTRINK-WWSNLPGYTIDSYKNSGMWSIGI 971

Search completed: March 10, 2004, 09:13:53
Job time : 7.2179 secs

GenCore version 5.1.6
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OM protein - protein search, using SW model

Run on: March 10, 2004 08:58:54 : Search time 12.0019 Seconds

(without alignments)

133.345 Million cell updates/sec

Title: US-09-848-834A.9

Perfect score: 160

Sequence: 1 KLISeIKGVIVHLEGVEGPSLHWSYGLRPX 31

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 389414 seqs, 51625971 residues

Total number of hits satisfying chosen parameters:

389414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
 Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents AA:^{*}
 1: /cgns2_6/ptodata/2/iaa/5A_COMBO.pep:
 2: /cgns2_6/ptodata/2/iaa/5B_COMBO.pep:
 3: /cgns2_6/ptodata/2/iaa/6A_COMBO.pep:
 4: /cgns2_6/ptodata/2/iaa/6B_COMBO.pep:
 5: /cgns2_6/ptodata/2/iaa/PCTUS_COMBO.pep:
 6: /cgns2_6/ptodata/2/iaa/backfile1.pep:
^{*}

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match	Length	DB ID	Description
1	117	73.1	42	1	US-09-446-692-20	Sequence 20, APP1
2	117	73.1	42	2	US-09-446-692-20	Sequence 19, APP1
3	116	72.5	27	1	US-09-446-692-19	Sequence 19, APP1
4	116	72.5	27	2	US-09-488-351A-19	Sequence 36, APP1
5	116	72.5	27	3	US-09-100-414B-36	Sequence 36, APP1
6	116	72.5	27	3	US-09-303-323-36	Sequence 36, APP1
7	116	72.5	27	4	US-09-70-014-36	Sequence 33, APP1
8	116	72.5	45	1	US-09-446-692-33	Sequence 33, APP1
9	116	72.5	45	2	US-09-488-351A-33	Sequence 43, APP1
10	113	70.6	27	3	US-09-100-414B-43	Sequence 43, APP1
11	113	70.6	27	3	US-09-303-323-43	Sequence 43, APP1
12	113	70.6	27	4	US-09-70-014-43	Sequence 55, APP1
13	68.8	31	3	US-09-100-414B-55	Sequence 55, APP1	
14	110	68.8	31	3	US-09-303-323-55	Sequence 55, APP1
15	110	68.8	31	4	US-09-70-014-55	Sequence 47, APP1
16	108	67.5	27	3	US-09-100-414B-47	Sequence 47, APP1
17	108	67.5	27	3	US-09-303-323-47	Sequence 47, APP1
18	108	67.5	27	3	US-09-303-323-47	Sequence 47, APP1
19	108	67.5	27	4	US-09-70-014-41	Sequence 47, APP1
20	108	67.5	27	4	US-09-70-014-47	Sequence 45, APP1
21	108	67.5	45	3	US-09-100-414B-45	Sequence 45, APP1
22	108	67.5	45	3	US-09-303-323-45	Sequence 45, APP1
23	108	67.5	45	4	US-09-70-014-45	Sequence 59, APP1
24	108	67.5	31	3	US-09-100-414B-59	Sequence 59, APP1
25	107	66.9	31	3	US-09-303-323-59	Sequence 59, APP1
26	107	66.9	31	4	US-09-70-014-59	Sequence 59, APP1
27	107	66.9	31	4	US-09-70-014-59	Sequence 59, APP1

RESULT 1
 US-09-446-692-20
 Sequence 20, Application US/08446692
 Patent No. 575951
 GENERAL INFORMATION:
 APPLICANT: Ladd, Anna
 APPLICANT: Wang, Chang Yi
 APPLICANT: Zamb, Timothy
 TITLE OF INVENTION: Immunogenic LHRH peptide constructs
 TITLE OF INVENTION: and synthetic universal immune stimulators for vaccines
 NUMBER OF SEQUENCES: 114
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Maria C.H. Lin
 STREET: 345 Park Avenue
 CITY: New York
 STATE: NY
 COUNTRY: US
 ZIP: 10154-0053
 COMPUTER READABLE COPY:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/446-692
 FILING DATE: 7-JUN-1995
 CLASSIFICATION: 424
 ATTORNEY/AGENT INFORMATION:
 NAME: Maria C.H. Lin
 REGISTRATION NUMBER: 29,323
 REFERENCE/DOCKET NUMBER: 1151-4146 US2
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (212)415-8745
 TELEFAX: (516)751-6849
 INFORMATION FOR SEQ ID NO: 20:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 42 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 MOLECULE TYPE: peptide
 US-09-446-692-20
 Query Match 73.1%; Score 117; DB 1; Length 42;
 Best Local Similarity 82.4%; Mismatches 2; Indels 2; Gaps 1;
 Matches 24;
 Qy 2 LLSIEIKGVIVRLLEGVEGPSLHWSYGLRP 30
 Db 15 VLSIEIKGVIVRLLEGVEGPSLHWSYGLRP 41

ALIGNMENTS

RESULT 2
 US-08-488-351A-20
 Sequence 20; Application US/08488351A
 Patent No. 5843146
 GENERAL INFORMATION:
 APPLICANT: Ladd, Anna
 APPLICANT: Wang, Chang Yi
 APPLICANT: Zamb, Timothy
 TITLE OF INVENTION: Immunogenic LHRH Peptide constructs
 NUMBER OF SEQUENCES: 114
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Maria C.H. Lin
 STREET: 345 Park Avenue
 CITY: New York
 STATE: NY
 ZIP: 10154-0053
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/488,351A
 FILING DATE: 7-JUN-1995
 CLASSIFICATION: 424
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/446,692
 FILING DATE: 7-JUN-1995
 CLASSIFICATION: 424
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/229,275
 FILING DATE: 14-APR-1994
 CLASSIFICATION: 424
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/057,166
 FILING DATE: 27-APR-1992
 CLASSIFICATION: 424
 ATTORNEY/AGENT INFORMATION:
 NAME: Maria C.H. Lin
 REGISTRATION NUMBER: 29,323
 REFERENCE/DOCKET NUMBER: 1151-4146 US2
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (212)415-8745
 TELEFAX: (516)751-6849
 SEQUENCE FOR SEQ ID NO: 20;
 SEQUENCE CHARACTERISTICS:
 LENGTH: 42 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 MOLECULE TYPE: peptide

US-08-488-351A-19
 Sequence 19; Application US/08488351A
 Patent No. 5843446
 GENERAL INFORMATION:
 APPLICANT: Ladd, Anna
 APPLICANT: Wang, Chang Yi
 APPLICANT: Zamb, Timothy
 TITLE OF INVENTION: Immunogenic LHRH Peptide constructs
 TITLE OF INVENTION: Immunoactive LHRH Peptide constructs
 NUMBER OF SEQUENCES: 114
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Maria C.H. Lin
 STREET: 345 Park Avenue
 CITY: New York
 STATE: NY
 ZIP: 10154-0053
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/488,351A
 FILING DATE: 7-JUN-1995
 CLASSIFICATION: 424
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/446,692
 FILING DATE: 7-JUN-1995
 CLASSIFICATION: 424
 TITLE OF INVENTION: Immunogenic LHRH Peptide constructs

RESULT 3
 US-08-488-351A-19
 Sequence 19; Application US/08488351A
 Patent No. 5753551
 GENERAL INFORMATION:
 APPLICANT: Ladd, Anna
 APPLICANT: Wang, Chang Yi
 APPLICANT: Zamb, Timothy
 TITLE OF INVENTION: Immunogenic LHRH Peptide constructs

PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/229,275
; FILING DATE: 14-APR-1994
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/057,166
; FILING DATE: 27-APR-1992
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: Mari C.H. Lin
; REGISTRATION NUMBER: 29,323
; REFERENCE/DOCKET NUMBER: 1151-4146 US2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212)415-8745
; TELEX/FAX: (516)751-6849
; INFORMATION FOR SEQ ID NO: 19:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-488-351A-19

Query Match 72.5%; Score 116; DB 2; Length 27;
Best Local Similarity 85.7%; Pred. No. 4e-11;
Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;

Qy 3 LSEIKGVIVHLRLEGVEGSPSLMSYGLRP 30
Db 1 LSEIKGVIVHLRLEGVGSE -HWSYGLRP 26

RESULT 6
US-09-303-323-36
; Sequence 36, Application US/09303323
; Patent No. 6228987
; GENERAL INFORMATION:
; APPLICANT: Wang, Chang Yi
; TITLE OF INVENTION: NOVEL LHRH PEPTIDE
; NUMBER OF SEQUENCES: 106
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Morgan & Finnegan, L.L.P.
; STREET: 345 Park Avenue
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10154-0054
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC Windows
; SOFTWARE: Word 97
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/303,323
; FILING DATE: 30-APR-1999
; CLASSIFICATION:
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: 10154-0054
; FILING DATE: 30-APR-1999
; ATTORNEY/AGENT INFORMATION:
; NAME: Maria H. Lin
; REGISTRATION NUMBER: 29,323
; REFERENCE/DOCKET NUMBER: 1151-4157
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 212-758-4800
; TELEX/FAX: 212-751-6849
; INFORMATION FOR SEQ ID NO: 36:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: Peptide
US-09-303-323-36

Query Match 72.5%; Score 116; DB 3; Length 27;
Best Local Similarity 85.7%; Pred. No. 4e-11;
Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;

Qy 3 LSEIKGVIVHLRLEGVEGSPSLMSYGLRP 30
Db 1 LSEIKGVIVHLRLEGVGGE -HWSYGLRP 26

RESULT 7
US-09-303-323-36
; Sequence 36, Application US/09303323
; Patent No. 6228987
; GENERAL INFORMATION:
; APPLICANT: Wang, Chang Yi
; TITLE OF INVENTION: NOVEL LHRH PEPTIDE
; NUMBER OF SEQUENCES: 106
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Morgan & Finnegan, L.L.P.
; STREET: 345 Park Avenue
; CITY: New York

Query Match 72.5%; Score 116; DB 3; Length 27;

STATE: NY USA
 COUNTRY: USA
 ZIP: 10154-0054
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC Windows
 SOFTWARE: Word 97
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/09/770,014
 FILING DATE:
 CLASSIFICATION:
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 09/100,414
 FILING DATE: 20-JUNE-1998
 ATTORNEY/AGENT INFORMATION:
 NAME: Maria H. Lin
 REGISTRATION NUMBER: 29,323
 REFERENCE/DOCKET NUMBER: 1151-4157
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 212-758-4800
 TELEFAX: 212-751-6849
 INFORMATION FOR SEQ ID NO: 36:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 27 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 MOLECULE TYPE: Peptide
 US-09-770-014-36

Query Match Score 116; DB 4; Length 27;
 Best Local Similarity 85.7%; Pred. No. 4e-11;
 Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;

Query Match Score 116; DB 1; Length 45;
 Best Local Similarity 85.7%; Pred. No. 7.3e-11;
 Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;

RESULT 9
 US-08-488-351A-33
 Sequence 33, Application US/08488351A
 Patent No. 5843446
 GENERAL INFORMATION:
 APPLICANT: Ladd, Anna
 APPLICANT: Wang, Chang Yi
 APPLICANT: Zamb, Timothy
 TITLE OF INVENTION: Immunogenic LHRH peptide constructs
 TITLE OF INVENTION: and synthetic universal immune stimulators for vaccines
 NUMBER OF SEQUENCES: 114
 CORRESPONDENCE ADDRESS:
 ADDRESS: Maria C.H. Lin
 STREET: 345 Park Avenue
 CITY: New York
 STATE: NY
 COUNTRY: US
 ZIP: 10154-0053
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: FacetIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/488,351A
 FILING DATE: 7-JUN-1995
 CLASSIFICATION: 424
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/229,275
 FILING DATE: 14-APR-1994
 CLASSIFICATION: 424
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/446,692
 FILING DATE: 7-JUN-1995
 CLASSIFICATION: 424
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/057,166
 FILING DATE: 27-APR-1992
 CLASSIFICATION: 424
 ATTORNEY/AGENT INFORMATION:
 NAME: Maria C.H. Lin
 REGISTRATION NUMBER: 29,323
 REFERENCE/DOCKET NUMBER: 1151-4146 US2
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (212)415-8745
 TELEFAX: (516)751-6849
 INFORMATION FOR SEQ ID NO: 33:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 45 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 MOLECULE TYPE: Peptide
 US-08-488-351A-33

Query Match Score 116; DB 2; Length 45;
 Best Local Similarity 85.7%;
 Pred. No. 7.3e-11;

/ RESULT 10
 / Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;
 / Sequence 43, Application US/09100414B
 / Pat. No. 6025468
 / GENERAL INFORMATION:
 / APPLICANT: Wang, Chang Yi
 / TITLE OF INVENTION: NOVEL LHRH PEPTIDE
 / NUMBER OF SEQUENCES: 106
 / CORRESPONDENCE ADDRESS:
 / ADDRESSEE: Morgan & Finnegan, L.L.P.
 / STREET: 345 Park Avenue
 / CITY: New York
 / STATE: NY
 / COUNTRY: USA
 / ZIP: 10154-0054
 / COMPUTER READABLE FORM:
 / MEDIUM TYPE: Floppy disk
 / COMPUTER: IBM PC compatible
 / OPERATING SYSTEM: PC Windows
 / SOFTWARE: Word 97
 / CURRENT APPLICATION DATA:
 / APPLICATION NUMBER: US/09/100,414B
 / FILING DATE: 20-JUNE-1998
 / CLASSIFICATION:
 / PRIORITY APPLICATION DATA:
 / PRIORITY NUMBER: 09/100,414
 / FILING DATE: 20-JUNE-1998
 / ATTORNEY/AGENT INFORMATION:
 / NAME: Maria H. Lin
 / REGISTRATION NUMBER: 29,323
 / REFERENCE/DOCKET NUMBER: 1151-4157
 / TELECOMMUNICATION INFORMATION:
 / TELEPHONE: 212-758-4800
 / TELEFAX: 212-751-6849
 / INFORMATION FOR SEQ ID NO: 43:
 / SEQUENCE CHARACTERISTICS:
 / LENGTH: 27 amino acids
 / TYPE: amino acid
 / TOPOLOGY: linear
 / MOLECULE TYPE: peptide
 / US-09-303-323-43
 / RESULT 11
 / Query Match 70.6%; Score 113; DB 3; Length 27;
 / Best Local Similarity 82.1%; Pred. No. 1; 2e-10;
 / Matches 23; Conservative 1; Mismatches 2; Indels 2; Gaps 1;
 / Sequence 43, Application US/090303323
 / Pat. No. 6228387
 / GENERAL INFORMATION:
 / APPLICANT: Wang, Chang Yi
 / TITLE OF INVENTION: NOVEL LHRH PEPTIDE
 / NUMBER OF SEQUENCES: 106
 / CORRESPONDENCE ADDRESS:
 / ADDRESSEE: Morgan & Finnegan, L.L.P.
 / STREET: 345 Park Avenue
 / CITY: New York
 / STATE: NY
 / COUNTRY: USA
 / ZIP: 10154-0054
 / COMPUTER READABLE FORM:
 / PRIORITY APPLICATION DATA:
 / PRIORITY NUMBER: 09/100,414
 / FILING DATE: 20-JUNE-1998
 / ATTORNEY/AGENT INFORMATION:
 / NAME: Maria H. Lin
 / REGISTRATION NUMBER: 29,323
 / REFERENCE/DOCKET NUMBER: 1151-4157
 / TELECOMMUNICATION INFORMATION:
 / TELEPHONE: 212-758-4800
 / TELEFAX: 212-751-6849
 / INFORMATION FOR SEQ ID NO: 43:
 / SEQUENCE CHARACTERISTICS:
 / LENGTH: 27 amino acids

TYPE: amino acid
 TOPOLOGY: linear
 MOLECULE TYPE: Peptide
 US-09-770-014-43

Query Match 70.6%; Score 113; DB 4; Length 27;
 Best Local Similarity 82.1%; P-Ped. No. 1.2e-10;
 Matches 23; Conservative 1; Mismatches 2; Indels 2; Gaps 1;

QY 3 LSEIKGVIVHLLEGEGVGLSLHWSYGLRP 30
 DB 1 LSEIKGVIVHLLEGVGGE - HWSYGLRP 26

RESULT 13

US-09-100-414B-55

Sequence 55, Application US/09100414B

Patent No. 6025468

GENERAL INFORMATION:

APPLICANT: Wang, Chang Yi

TITLE OF INVENTION: NOVEL LHRH PEPTIDE

NUMBER OF SEQUENCES: IMMUNOGENS

106

CORRESPONDENCE ADDRESS:

ADDRESSEE: Morgan & Finnegan, L.L.P.

STREET: 345 Park Avenue

CITY: New York

STATE: NY

COUNTRY: USA

ZIP: 10154-0054

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC Windows

SOFTWARE: Word 97

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/303,323

FILING DATE: 30-APR-1999

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 09/100,414

FILING DATE: 20-JUNE-1998

ATTORNEY/AGENT INFORMATION:

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 55:

SEQUENCE CHARACTERISTICS:

LENGTH: 31 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 55:

SEQUENCE CHARACTERISTICS:

LENGTH: 31 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 30:

SEQUENCE CHARACTERISTICS:

LENGTH: 30 amino acids

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-100-414B-55

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 11151-4157

TELECOMMUNICATION INFORMATION:

/ REFERENCE/DOCKET NUMBER: 1151-4157
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 212-751-4800
/ TELFAX: 212-751-849
/ INFORMATION FOR SEQ ID NO: 55:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 31 amino acids
/ TYPE: amino acid
/ TOPOLOGY: linear
/ MOLECULE TYPE: peptide
us-09-770-014-55

Query Match 68.8%; Score 110; DB 4; Length 31;
Best Local Similarity 78.6%; Pred. No. 4e-10;
Matches 22; Conservative 1; Mismatches 5; Indels 0; Gaps 0;
Qy 3 LSEIKGTVHLEGVCGPSLMSYGLRP 30
Db 3 LSEIKGTVHLEGVCGPSLMSYGLRP 30

Search completed: March 10, 2004, 09:28:53
Job time : 12.0019 secs

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OM protein - protein search, using SW model.

Run on: March 10, 2004, 09:16:59 ; Search time 24.3658 Seconds

(without alignment)

268 645 Million cell updates/sec

Title: US-09-848-834A-9

Perfect score: 160

Sequence: 1 KLISEIKGVIVTHLEGVEGFSLHWSYGLRPX 31

Scoring table: BLOSUM62

Gapopen 10.0 , Gapext 0.5

Searched: 809742 seqs, 211153259 residues

Total number of hits satisfying chosen parameters:

809742

Post-processing: Minimum Match 100%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA: *

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1: /cgm2_6/prodata/2/pubpas/us07_pubcomb.pep:*
2: /cgm2_6/prodata/2/pubpas/pct_new_pub.pep:*
3: /cgm2_6/prodata/2/pubpas/us05_new_pub.pep:*
4: /cgm2_6/prodata/2/pubpas/us06_pubcomb.pep:*
5: /cgm2_6/prodata/2/pubpas/us03_pubcomb.pep:*
6: /cgm2_6/prodata/2/pubpas/pctns_pubns.pep:*
7: /cgm2_6/prodata/2/pubpas/us08_new_pub.pep:*
8: /cgm2_6/prodata/2/pubpas/us09_pubcomb.pep:*
9: /cgm2_6/prodata/2/pubpas/us09a_pubcomb.pep:*
10: /cgm2_6/prodata/2/pubpas/us09b_pubcomb.pep:*
11: /cgm2_6/prodata/2/pubpas/us09c_pubcomb.pep:*
12: /cgm2_6/prodata/2/pubpas/us09_new_pub.pep:*
13: /cgm2_6/prodata/2/pubpas/us10_pubcomb.pep:*
14: /cgm2_6/prodata/2/pubpas/us10b_pubcomb.pep:*
15: /cgm2_6/prodata/2/pubpas/us10c_pubcomb.pep:*
16: /cgm2_6/prodata/2/pubpas/us10_new_pub.pep:*
17: /cgm2_6/prodata/2/pubpas/us10_new_pub.pep:*
18: /cgm2_6/prodata/2/pubpas/us10_pubcomb.pep:*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length DB ID	Description
1	159	99.4	Sequence 9, Appl
2	159	99.4	Sequence 9, Appl
3	116	72.5	Sequence 9, Appl
4	116	72.5	Sequence 9, Appl
5	99	61.9	Sequence 9, Appl
6	99	61.9	Sequence 9, Appl
7	86	53.8	Sequence 9, Appl
8	79	49.4	Sequence 9, Appl
9	79	49.4	Sequence 9, Appl
10	78	48.8	Sequence 9, Appl
11	78	48.8	Sequence 9, Appl
12	78	48.8	Sequence 9, Appl
13	77	48.1	Sequence 9, Appl
14	77	48.1	Sequence 9, Appl
15	73	45.6	Sequence 9, Appl

Sequence 12, Appl
Sequence 11, Appl
Sequence 19, Appl
Sequence 20, Appl
Sequence 16, Appl
Sequence 30, Appl
Sequence B, Appl
Sequence 22, Appl
Sequence 20, Appl
Sequence 10, Appl
Sequence 50, Appl
Sequence 18, Appl
Sequence 20, Appl
Sequence 29, Appl
Sequence 3, Appl
Sequence 48, Appl
Sequence 40, Appl
Sequence 51, Appl
Sequence 14, Appl
Sequence 35, Appl
Sequence 38, Appl
Sequence 40, Appl
Sequence 42, Appl
Sequence 51, Appl
Sequence 13, Appl
Sequence 14, Appl
Sequence 35, Appl
Sequence 38, Appl
Sequence 40, Appl
Sequence 42, Appl
Sequence 27, Appl
Sequence 30, Appl
Sequence 32, Appl
Sequence 34, Appl
Sequence 49, Appl

ALIGNMENTS

RESULT 1
US-09-848-834A-9
Sequence 9, Application US/09848834A
Patent No. US2002007616A1
GENERAL INFORMATION:
APPLICANT: Ashton Corporation
TITLE OF INVENTION: Chimeric Peptide Immunogens
FILE REFERENCE: 1102865-0047
CURRENT APPLICATION NUMBER: US/09848-834A
CURRENT FILING DATE: 2001-05-04
PRIOR APPLICATION NUMBER: 60/202,328
PRIOR FILING DATE: 2000-05-05
NUMBER OF SEQ ID NOS: 20
SOFTWARE: PatentIn version 3.0
SEQ ID NO 9
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Chimeric peptide made up of amino acid sequence 288-302 of the measles virus fusion protein, F linked by a spacer peptide to amino acid sequence 2-10 of the GnRH hormone

; OTHER INFORMATION: Amidated glycine or glycaminide
US-09-848-834a-9

Query Match 99.4%; Score 159; DB 9; Length 31;
Best Local Similarity 100.0%; Pred. No. 5.8e-16;
Matches 30; Conservative 0; Mismatches 0; Indels 0;
Gaps 0;
SEQ ID NO: 9

Qy 1 KLLSEIKGVIVHLLEGEGPSLHWSYGLRP 30
Db 1 KLLSEIKGVIVHLLEGEGPSLHWSYGLRP 30

RESULT 2
US-09-848-834a-17
Sequence 17, Application US/09848834A
Patent No. US20020076416A1
GENERAL INFORMATION:
APPLICANT: Aphton Corporation
TITLE OF INVENTION: Chimeric Peptide Immunogens
FILE REFERENCE: 1102865-0047
CURRENT FILING DATE: 2001-05-04
PRIOR APPLICATION NUMBER: US/09/848, 834A
NUMBER OF SEQ ID NOS: 20
SEQ ID NO: 17
LENGTH: 47
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Chimeric peptide consisting of amino acid sequence 1-10 of the GnRH hormone linked by a spacer to amino acid sequence 288-302 of the Measles virus protein F linked by a spacer to amino acid sequence 2-10 of the GnRH hormone
NAME/KEY: MOD_RES
LOCATION: (1) .. (1)
OTHER INFORMATION: Pyroglutamic acid or 5-Oxoprolidine
NAME/KEY: MOD_RES
LOCATION: (47) .. (47)
OTHER INFORMATION: Amidated-glycine or glycaminide
NAME/KEY: PEPTIDE
LOCATION: (1) .. (10)
OTHER INFORMATION: Amino acid sequence 1-10 of the human GnRH hormone
NAME/KEY: PEPTIDE
LOCATION: (11) .. (18)
OTHER INFORMATION: Spacer peptide
NAME/KEY: PEPTIDE
LOCATION: (19) .. (34)
OTHER INFORMATION: Amino acid sequence 288-302 of the Measles virus fusion protein,
NAME/KEY: PEPTIDE
LOCATION: (35) .. (38)
OTHER INFORMATION: Spacer peptide
NAME/KEY: PEPTIDE
LOCATION: (39) .. (47)

OTHER INFORMATION: Amino acid sequence 2-10 of the human GnRH hormone
US-09-848-834a-17
Query Match 99.4%; Score 159; DB 9; Length 47;
Best Local Similarity 100.0%; Pred. No. 9.2e-16;
Matches 30; Conservative 0; Mismatches 0; Indels 0;
Gaps 0;
SEQ ID NO: 9

Qy 1 KLLSEIKGVIVHLLEGEGPSLHWSYGLRP 30
Db 17 KLLSEIKGVIVHLLEGEGPSLHWSYGLRP 46

RESULT 3
US-10-076-674-9
Sequence 9, Application US/10076674
Publication No. US20030165478A1
GENERAL INFORMATION:
APPLICANT: Sokoll, Kenneth K.
; FEATURE:

; TITLE OF INVENTION: Stabilized Synthetic Immunogen Delivery System
; FILE REFERENCE: Immunogen Delivery System
; CURRENT APPLICATION NUMBER: US/10/076,674
; CURRENT FILING DATE: 2002-04-23
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 9
; LENGTH: 45
; TYPE: PRT
; ORGANISM: Human
; US-10-076-674-9
; Query Match 72.5%; Score 116; DB 14; Length 45;
; Best Local Similarity 85.7%; Pred. No. 1.5e-09;
; Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;
; Qy 3 LSEIKGVIVHLLEGEGPSLHWSYGLRP 30
; Db 19 LSEIKGVIVHLLEGEGGE-HWSYGLRP 44

RESULT 4
US-10-355-161A-9
; Sequence 9, Application US/10355161A
; Publication No. US200400003897A1
; GENERAL INFORMATION:
; APPLICANT: Sokoll, Kenneth K.
; TITLE OF INVENTION: Stabilized Synthetic Immunogen Delivery System
; FILE REFERENCE: Immunogen Delivery System
; CURRENT APPLICATION NUMBER: US/10/355,161A
; CURRENT FILING DATE: 2003-01-31
; PRIOR APPLICATION NUMBER: US/10/355,161A
; PRIOR FILING DATE: 2002-02-14
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 9
; LENGTH: 45
; TYPE: PRT
; ORGANISM: Human
; US-10-355-161A-9
; Query Match 72.5%; Score 116; DB 15; Length 45;
; Best Local Similarity 85.7%; Pred. No. 1.5e-09;
; Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;
; Qy 3 LSEIKGVIVHLLEGEGPSLHWSYGLRP 30
; Db 19 LSEIKGVIVHLLEGEGGE-HWSYGLRP 44

RESULT 5
US-09-847-102A-33
; Sequence 33, Application US/09847102A
; Publication No. US20030044409A1
; GENERAL INFORMATION:
; APPLICANT: University of California
; APPLICANT: Carlson, Maripat
; APPLICANT: Rhee, Chae-Seo
; APPLICANT: Loranzo, Leoni M.
; APPLICANT: Maloni, Sen
; TITLE OF INVENTION: IMMUNOLOGIC COMPOSITIONS AND METHODS FOR
; FILE REFERENCE: 2200-20639.00
; CURRENT APPLICATION NUMBER: US/09/847,102A
; CURRENT FILING DATE: 2001-05-01
; NUMBER OF SEQ ID NOS: 138
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO: 33
; LENGTH: 75
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:

OTHER INFORMATION: Chimeric peptide consisting of amino acid sequence 1-10 of the US-09-848-102A-33
 OTHER INFORMATION: RH hormone linked by a spacer to amino acid sequence 288-302 of the Measles virus fusion protein,
 NAME/KEY: PEPTIDE
 LOCATION: (1) .(10)
 OTHER INFORMATION: Amino acid sequence 1-10 of the human GnRH hormone
 NAME/KEY: PEPTIDE
 LOCATION: (11) .(18)
 OTHER INFORMATION: Spacer peptide
 NAME/KEY: PEPTIDE
 LOCATION: (19) .(34)
 OTHER INFORMATION: Amino acid sequence 288-302 of the Measles
 OTHER INFORMATION: Virus fusion protein, F
 NAME/KEY: MOD-RES
 LOCATION: (1) .(1)
 OTHER INFORMATION: Pyroglutamic acid or 5-oxoproline
 US-09-848-834A-13

RESULT 6
 US-10-285-976-231
 Sequence 231; Application US/10285976
 Publication No. US2003165500A1
 GENERAL INFORMATION:
 APPLICANT: Rhee, Chae-Seo
 APPLICANT: Mallin, Sean
 APPLICANT: Wu, Christina
 APPLICANT: Leoni, Lorenzo M.
 APPLICANT: Corr, Maripat
 APPLICANT: Carson, Dennis A.
 APPLICANT: The Regents of the University of California
 TITLE OF INVENTION: Wnt and Frizzled Receptors as Targets for Immunotherapy
 FILE REFERENCE: 023070-130320US
 CURRENT APPLICATION NUMBER: US/10/285,976
 CURRENT FILING DATE: 2002-11-01
 PRIOR APPLICATION NUMBER: US 60/287,995
 PRIOR FILING DATE: 2001-05-01
 PRIOR APPLICATION NUMBER: WO PCT/US02/13602
 PRIOR FILING DATE: 2002-05-01
 NUMBER OF SEQ ID NOS: 232
 SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO: 231
 LENGTH: 75
 TYPE: PRT
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: Description of Artificial Sequence: PMMVF-ZD2
 OTHER INFORMATION: measles virus fusion (MVF) epitope fused to
 OTHER INFORMATION: frizzled domain
 US-10-285-976-231

Query Match 61.9%; Score 99; DB 14; Length 75;
 Best Local Similarity 95.5%; Pred. No. 7.7e-07; Indels 0; Gaps 0;
 Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 KLLSEIKGVIVHLRLEGVGPSL 22
 Db 2 KLLSLIKGVIVHLRLEGVGPSL 23

RESULT 7
 US-09-848-834A-13
 Sequence 13; Application US/09848834A
 Patent No. US200202076416A1
 GENERAL INFORMATION:
 APPLICANT: Abidion Corporation
 TITLE OF INVENTION: Chimeric Peptide Immunogens
 FILE REFERENCE: 1102865-0047
 CURRENT APPLICATION NUMBER: US/09/848,834A
 CURRENT FILING DATE: 2001-15-04
 PRIOR APPLICATION NUMBER: 60/202,328
 PRIOR FILING DATE: 2000-05-05
 NUMBER OF SEQ ID NOS: 20
 SOFTWARE: PatentIn version 3.0
 SEQ ID NO: 10
 LENGTH: 34
 TYPE: PRT
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: Tentoxylisin
 OTHER INFORMATION: Tetanus toxoid precursor (Tentoxysin) linked by a spacer to a
 OTHER INFORMATION: ino acid sequence 2-10 of the GnRH hormone
 NAME/KEY: MOD-RES
 LOCATION: (1) .(1)
 OTHER INFORMATION: Amidated phenylalanine
 NAME/KEY: PEPTIDE
 LOCATION: (1) .(21)
 OTHER INFORMATION: Amino acids 947-967 of the Tetanus Toxoid Precursor
 OTHER INFORMATION: (Tentoxylisin)
 NAME/KEY: PEPTIDE
 LOCATION: (22) .(25)
 OTHER INFORMATION: Spacer peptide
 NAME/KEY: PEPTIDE
 LOCATION: (26) .(34)
 OTHER INFORMATION: Amino acids 2-10 of the human GnRH hormone
 NAME/KEY: MOD-RES
 LOCATION: (34) .(34)
 OTHER INFORMATION: Amidated glycine or glycaminamide
 US-09-848-834A-10

Query Match 49.4%; Score 79; DB 9; Length 34;
 Best Local Similarity 92.9%; Pred. No. 0.0025; Indels 0; Gaps 0;
 Matches 13; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 VEGPSLHNSYGLRP 30
 Db 2 :|||||||

Db 20 LEGPSLHWSYGLRP 33
RESULT 9
US-09-848-834A-18
Sequence 19, Application US/09848834A
Patent No US200006416A1
GENERAL INFORMATION
APPLICANT: Aphton Corporation
TITLE OF INVENTION: Chimeric Peptide Immunogens
FILE REFERENCE: 1102865-0047
CURRENT APPLICATION NUMBER: US/09/848,834A
CURRENT FILING DATE: 2001-05-04
PRIOR APPLICATION NUMBER: 60/202,328
PRIOR FILING DATE: 2000-05-05
NUMBER OF SEQ ID NOS: 20
SOFTWARE: PatentIn version 3.0
SEQ ID NO 18
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Chimeric peptide consisting of amino acid sequence 1-10 of human
OTHER INFORMATION: GnRH linked by a spacer to amino acid sequence 947-967 of the Tet
OTHER INFORMATION: anus toxoid precursor (tentoxin) Protein Linked by a spacer to
OTHER INFORMATION: o amino acid sequence 2-10 of human GnRH
NAME/KEY: MOD_RES
LOCATION: (1)..(1)
OTHER INFORMATION: Amino acid sequence 1-10 of the human GnRH hormone
NAME/KEY: MOD_RES
LOCATION: (50)..(50)
OTHER INFORMATION: Amidated glycine or glycaminide
NAME/KEY: PEPTIDE
LOCATION: (17)..(37)
OTHER INFORMATION: Amino acid sequence 947-967 of the Tetanus toxoid precursor (Tet
NAME/KEY: PEPTIDE
LOCATION: (11)..(16)
OTHER INFORMATION: Spacer peptide
NAME/KEY: PEPTIDE
LOCATION: (42)..(50)
OTHER INFORMATION: Amino acid sequence 2-10 of the human GnRH hormone
US-09-848-834A-18
Query Match 49.4%; Score 79; DB 9; Length 50;
Best Local Similarity 99.9%; Pred. No. 0.00039;
Matches 13; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
Qy 17 VEGPSLHWSYGLRP 30
Db 36 LEGPSLHWSYGLRP 49

Db 21 LEGPSLHWSYGLRP 33
RESULT 10
US-10-223-711-10
Sequence 10, Application US/10223711
Publication No. US20030113344A1
GENERAL INFORMATION
APPLICANT: Bakalecz, Lauren O.
APPLICANT: Kaumaya, Pravin T.P.
TITLE OF INVENTION: Synthetic Chimeric Fimbrin Peptides
FILE REFERENCE: 11525/04059
CURRENT APPLICATION NUMBER: US/10/223,711
CURRENT FILING DATE: 2003-08-19
PRIOR APPLICATION NUMBER: 09/148,711
PRIOR FILING DATE: 1998-09-04
PRIOR APPLICATION NUMBER: 08/460,502

Db 22 LEGPSLHWSYGLRP 33
RESULT 11
US-09-847-102A-31
Sequence 31, Application US/09847102A
Publication No. US2003004409A1
GENERAL INFORMATION
APPLICANT: University of California
APPLICANT: Carlson, Dennis A.
APPLICANT: Corr, Maripat
APPLICANT: Rhee, Chae-Soo
APPLICANT: Lorenzo, Leoni M.
APPLICANT: Malini, Sen
APPLICANT: Title of Invention: IMMUNOLOGIC COMPOSITIONS AND METHODS FOR
OTHER INFORMATION: STUDYING AND TREATING CANCERS EXPRESSING FRIZZLED ANTIGENS
APPLICANT: Title of Invention: STUDYING AND TREATING CANCERS EXPRESSING FRIZZLED ANTIGENS
APPLICANT: File Reference: 22000-20629,00
APPLICANT: Current Application Number: US/09/847,102A
APPLICANT: Current Filing Date: 2001-05-01
APPLICANT: Number of Seq ID Nos: 138
APPLICANT: Software: PastSeq for Windows Version 4.0
SEQ ID NO 31
LENGTH: 75
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: PFZD2-MMVF
US-09-847-102A-31
Query Match 48.8%; Score 78; DB 10; Length 75;
Best Local Similarity 94.4%; Pred. No. 0.00085;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 KLLSLIKGVIVHLRLEGVE 18
Db 58 KLLSLIKGVIVHLRLEGVE 75

Db 23 LEGPSLHWSYGLRP 40
RESULT 12
US-10-285-976-229
Sequence 229, Application US/10285976
Publication No. US2003016550A1
GENERAL INFORMATION
APPLICANT: Wu, Christina
APPLICANT: Malini, Sen
APPLICANT: Rhee, Chae-Soo
APPLICANT: Leon, Lorenzo M.
APPLICANT: Corr, Maripat
APPLICANT: Carlson, Dennis A.
APPLICANT: The Regents of the University of California
TITLE OF INVENTION: Wnt and Frizzled Receptors as Targets for Immunotherapy
APPLICANT: Title of Invention: in Head and Neck Squamous Cell Carcinomas
FILE REFERENCE: 02307-O-13032005
CURRENT APPLICATION NUMBER: US/10/285,976
CURRENT FILING DATE: 2002-11-01
PRIORITY APPLICATION NUMBER: US 60/287,995

PRIOR FILING DATE: 2001-05-01
 PRIORITY APPLICATION NUMBER: WO PCT/US02/13802
 PRIOR FILING DATE: 2002-05-01
 NUMBER OF SEQ ID NOS: 232
 SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO: 229
 LENGTH: 75
 TYPE: PRT
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: Description of Artificial Sequence: PFZD2 - MNVF
 OTHER INFORMATION: measles virus fusion (MVF) epitope fused to
 OTHER INFORMATION: measles virus fusion protein
 OTHER INFORMATION: frizzled domain
 US-10-285-976-229

Query Match Score 78; DB 14; Length 75;
 Best Local Similarity 94.4%; Pred. No. 0.00085; Mismatches 1; Indels 0; Gaps 0;
 Matches 17; Conservative 0; Gaps 0;

Oy 1 LSEIKGVIVHLRLEGVE 18
 Db 58 KULLSLKGIVVHLRLEGVE 75

RESULT 13
 US-09-848-834A-8
 Sequence 8, Application US/09848834A
 Patent No. US20020076416A1
 GENERAL INFORMATION:
 APPLICANT: Adhton Corporation
 TITLE OF INVENTION: Chimeric Peptide Immunogens
 FILE REFERENCE: 1102865-0047
 CURRENT APPLICATION NUMBER: US/09/848,834A
 CURRENT FILING DATE: 2001-05-04
 PRIOR APPLICATION NUMBER: 60/202,328
 PRIOR FILING DATE: 2000-05-05
 NUMBER OF SEQ ID NOS: 20
 SOFTWARE: PatentIn version 3.0
 SEQ ID NO: 8
 LENGTH: 16
 TYPE: PRT
 ORGANISM: Measles virus
 FEATURE:
 NAME/KEY: PEPTIDE
 LOCATION: (1).-(16)
 OTHER INFORMATION: Amino acid sequence 288-302 of the measles
 OTHER INFORMATION: Amino acid fusion protein, F

US-09-848-834A-8

Query Match Score 77; DB 9; Length 16;
 Best Local Similarity 100.0%; Pred. No. 0.00021; Mismatches 0; Indels 0; Gaps 0;
 Matches 16; Conservative 0; Gaps 0;

Oy 3 LSEIKGVIVHLRLEGVE 18
 Db 1 LSEIKGVIVHLRLEGVE 16

RESULT 14
 US-10-411-544-32
 Sequence 32, Application US/10411544
 Publication No. US2003232758A1
 GENERAL INFORMATION:
 APPLICANT: St. George-Hyslop, Peter
 APPLICANT: McLaurin, Joanne
 TITLE OF INVENTION: Immunological Methods and Compositions for the Treatment of Alzheimer's Disease
 FILE REFERENCE: LI01547
 CURRENT APPLICATION NUMBER: US/10/411,544
 CURRENT FILING DATE: 2003-04-10
 NUMBER OF SEQ ID NOS: 52
 SOFTWARE: PatentIn version 3.1
 SEQ ID NO: 32

; LENGTH: 25
; TYPE: PRT
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: chimeric sequence
; US-10-411-544-32
; Query Match Score 77; DB 15; Length 25;
; Best Local Similarity 72.0%; Pred. No. 0.00035; Mismatches 7; Indels 0; Gaps 0;
; Matches 18; Conservative 0; Gaps 0;
; Qy 3 LSEIKGVIVHLRLEGVEPSLWISYI 27
; Db 1 LSEIKGVIVHLRLEGGSFRHDSGYG 25
; RESULT 15
; US-09-848-834A-11
; Sequence 11, Application US/09848834A
; Patent No. US20020076416A1
; GENERAL INFORMATION:
; APPLICANT: Adhton Corporation
; TITLE OF INVENTION: Chimeric Peptide Immunogens
; FILE REFERENCE: 1102865-0047
; CURRENT APPLICATION NUMBER: US/09/848,834A
; CURRENT FILING DATE: 2001-05-04
; PRIOR APPLICATION NUMBER: 60/202,328
; PRIOR FILING DATE: 2000-05-05
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO: 11
; LENGTH: 28
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Chimeric peptide consisting of amino acid sequence 830-844 of tetanus toxoid precursor (Tentoxylisin) linked by a spacer to a
; OTHER INFORMATION: Tetanus toxoid precursor (Tentoxylisin) linked by a spacer to a
; NAME/KEY: MOD-RES
; LOCATION: (1).-(1)
; OTHER INFORMATION: Amidated-glycine
; NAME/KEY: MOD-RES
; LOCATION: (28).-(28)
; OTHER INFORMATION: Amidated-glycine or glycaminide
; NAME/KEY: PEPTIDE
; LOCATION: (1).-(15)
; OTHER INFORMATION: Amino acid sequence 830-844 of the tetanus Toxoid Precursor
; NAME/KEY: PEPTIDE
; LOCATION: (16).-(19)
; OTHER INFORMATION: Spacer peptide
; NAME/KEY: PEPTIDE
; LOCATION: (20).-(28)
; OTHER INFORMATION: Amino acid sequence 2-10 of the human GnRH hormone
; NAME/KEY: PEPTIDE
; LOCATION: (16).-(19)
; OTHER INFORMATION: 100.0%; Pred. No. 0.0015; Mismatches 0; Indels 0; Gaps 0;

Query Match Score 73; DB 9; Length 28;
 Best Local Similarity 100.0%; Pred. No. 0.0015; Mismatches 0; Indels 0; Gaps 0;
 Matches 12; Conservative 0; Gaps 0;
 Qy 19 GPSLHWSYGLRP 30
 Db 16 GPSLHWSYGLRP 27
; Search completed: March 10, 2004, 10:25:48
; Job time : 25.3658 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: March 10, 2004, 08:58:54 : Search time 9.64981 Seconds

(without alignments) 309.015 Million cell updates/sec

Title: US-09-848-834A-9.

Perfect score: 160

Sequence: K1KSE1KGVTIVRLEGVEGPSLHWSYGLRPX 31

scoring table: BLOSUM62

Gapopen 10.0 , Gapext 0.5

Searched: 283366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : PIR78:*

1: Pir1:*

2: Pir2:*

3: Pir3:*

4: Pir4:*

Pred. No. is the number of results predicted by chance to have a a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match	Length	DB ID	Description
1	72	45.0	282	2	P00376	cell fusion glycop
2	72	45.0	282	2	PQ0388	cell fusion glycop
3	72	45.0	534	1	JT0274	cell fusion glycop
4	72	45.0	546	2	S47300	gene F protein - r
5	72	45.0	550	1	BA8556	cell fusion glycop
6	72	45.0	553	1	VGNZMV	cell fusion glycop
7	71	44.4	546	1	VGNZRK	cell fusion glycop
8	71	44.4	546	2	S47305	gene F protein - r
9	66.5	41.6	552	2	S47034	cell fusion glycop
10	66	41.2	546	1	VGNZR	cell fusion protein
11	65	40.6	542	2	JG2223	cell fusion protein
12	65	40.6	662	1	VGNZCD	cell fusion protein
13	65	40.6	662	2	S21382	cell fusion protein
14	64	40.0	631	1	VGNZPD	cell fusion glycop
15	64	40.0	631	1	A48346	cell fusion protein
16	60	37.5	546	2	S55386	gonadoliberin prec
17	57.5	35.9	92	1	RHTUG	gonadoliberin prec
18	57	35.6	90	1	REMSG	gonadoliberin prec
19	57	35.6	92	1	RHTG	gene F protein - r
20	56	35.0	636	2	S47299	gonadoliberin prec
21	54	33.8	67	2	I78541	spike glycoprotein
22	53	33.1	508	1	VGDNFR	gonadoliberin - pi
23	52	32.5	10	1	RHFGG	gonadoliberin - sh
24	52	32.5	10	2	I51423	phosphoribosylamin
25	52	32.5	89	1	DCBSPK	probable tetR-fami
26	52	32.5	379	1	T37168	hypothetical prote
27	50.5	31.6	333	2	T23151	conserved hypothet
28	50	31.2	451	2	AH0063	

ALIGNMENTS

RESULT 1
 PQ0376
 cell fusion Glycoprotein - measles virus (strain TR) (fragment)
 C;Species: measles virus
 C;Date: 17-Apr-1993 #Sequence_revision 17-Apr-1993
 C;Accession: PQ0376
 R;Schulz, T.F.; Head, J.G.; Whitby, D.; Tizard, E.J.; Dillon, M.J.; Weiss, R.A.
 J;Gen. Virol. 73, 1581-1586, 1992
 A;Title: A measles virus isolate from a child with Kawasaki disease: sequence comparison
 A;Reference number: PQ0374; PMID:92300360; PMID:1607874
 A;Accession: P00376
 A;Molecule type: genomic RNA
 C;Residues: 1-282 <SCH>
 C;Genetics:
 A;Gene: F
 C;Superfamily: paramyovirus cell fusion protein
 C;Keywords: glycoprotein; membrane fusion

Query Match %
 Best Local Similarity 100.0%; Pred. No. 0.016;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 1
 PQ0376
 cell fusion Glycoprotein - measles virus (strain Schwarz vaccine) (fragment)
 C;Species: measles virus
 C;Date: 17-Apr-1993 #Sequence_revision 17-Apr-1993
 C;Accession: PQ0378
 R;Schulz, T.F.; Head, J.G.; Whitby, D.; Tizard, E.J.; Dillon, M.J.; Weiss, R.A.
 J;Gen. Virol. 73, 1581-1586, 1992
 A;Title: A measles virus isolate from a child with Kawasaki disease: sequence comparison
 A;Reference number: PQ0374; PMID:92300360; PMID:1607874
 A;Accession: P00376
 A;Molecule type: genomic RNA
 C;Residues: 1-282 <SCH>
 C;Genetics:
 A;Gene: F
 C;Superfamily: paramyovirus cell fusion protein
 C;Keywords: glycoprotein; membrane fusion

Query Match %
 Best Local Similarity 100.0%; Pred. No. 0.016;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 2
 PQ0388
 cell fusion Glycoprotein - measles virus (strain Schwarz vaccine) (fragment)
 C;Species: measles virus
 C;Date: 17-Apr-1993 #Sequence_revision 17-Apr-1993
 C;Accession: PQ0388
 R;Schulz, T.F.; Head, J.G.; Whitby, D.; Tizard, E.J.; Dillon, M.J.; Weiss, R.A.
 J;Gen. Virol. 73, 1581-1586, 1992
 A;Title: A measles virus isolate from a child with Kawasaki disease: sequence comparison
 A;Reference number: PQ0374; PMID:92300360; PMID:1607874
 A;Accession: P00376
 A;Molecule type: genomic RNA
 C;Residues: 1-282 <SCH>
 C;Genetics:
 A;Gene: F
 C;Superfamily: paramyovirus cell fusion protein
 C;Keywords: glycoprotein; membrane fusion

Query Match %
 Best Local Similarity 100.0%; Pred. No. 0.016;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 2
 PQ0388
 cell fusion Glycoprotein - measles virus (strain Schwarz vaccine) (fragment)
 C;Species: measles virus
 C;Date: 17-Apr-1993 #Sequence_revision 17-Apr-1993
 C;Accession: PQ0388
 R;Schulz, T.F.; Head, J.G.; Whitby, D.; Tizard, E.J.; Dillon, M.J.; Weiss, R.A.
 J;Gen. Virol. 73, 1581-1586, 1992
 A;Title: A measles virus isolate from a child with Kawasaki disease: sequence comparison
 A;Reference number: PQ0374; PMID:92300360; PMID:1607874
 A;Accession: P00376
 A;Molecule type: genomic RNA
 C;Residues: 1-282 <SCH>
 C;Genetics:
 A;Gene: F
 C;Superfamily: paramyovirus cell fusion protein
 C;Keywords: glycoprotein; membrane fusion

Query Match %
 Best Local Similarity 100.0%; Pred. No. 0.016;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 3

JU0274 cell fusion glycoprotein precursor - subacute sclerosing panencephalitis virus (strain Y) cell fusion glycoprotein F1; fusion glycoprotein F1; feline glycoprotein F1; panencephalitis virus, SSPEV C;Species: subacute sclerosing panencephalitis virus, SSPEV C;Date: 31-Dec-1993 #text_change 16-Jun-2000 C;Accession: JU0274 R;Romase, K.; Haga, T.; Yoshikawa, Y.; Sato, T.A.; Yamanouchi, K. Virus Genes 4, 173-181, 1990 A;Title: Molecular analysis of structural protein genes of the Yamagata-1 strain of defective virus number: JU0274; MUID:90385702; PMID:1698327 A;Accession: JU0274 A;Molecule type: mRNA A;Residues: 1-534 <KOM> A;Cross-references: EMBL:D10548; NID:9222256; PIDN:BAA01405.1; PID:9222257 A;Note: the authors translated the codon GTA for residue 459 as Gly and GGG for residue 460 C;Genetics: A;Gene: F C;Superfamily: parainfluenza virus cell fusion protein C;Keywords: glycoprotein; membrane fusion; transmembrane protein C;KeyWords: membrane fusion protein; transmembrane protein C;1-22/Domain: signal sequence #status predicted <SG> F;23-107/Product: cell fusion glycoprotein F2 #status predicted <FF2> F;108-550/Product: cell fusion glycoprotein F2 #status predicted <FF2> F;108-550/Domain: hydrophobic transmembrane #status predicted <TM> F;6,29,61,67/Binding site: carbohydrate (Asn) (covalent) #status predicted <FF1> F;49-514/Domain: transmembrane #status predicted <TM> F;6,29,61,67/Binding site: carbohydrate (Asn) (covalent) #status predicted <FF1> Query Match Score 72; DB 1; Length 550; Best Local Similarity 100.0%; Pred. No. 0.033; Mismatches 0; Indels 0; Gaps 0; Matches 15; Conservative 15; Query 3 LSEIKGVIVHLREGV 17 Db 288 LSEIKGVIVHLREGV 302

RESULT 4

S4700 gene F protein - rinderpest virus C;Species: rinderpest virus C;Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 15-Oct-1999 C;Accession: S4700; P00865 R;Evans, S.A.; Baron, M.D.; Chamberlain, R.W.; Goatley, L.; Barrett, T. Submitted to the EMBL Data Bank, March 1994 A;Description: The complete nucleotide sequence of the fusion protein gene of the vaccinia virus A;Reference number: S4700 A;Accession: S4700 A;Molecule type: DNA A;Residues: 1-546 <EVA> A;Cross-references: EMBL:231656; NID:9535406; PIDN:CAA83482.1; PID:9535407 R;Chamberlain, R.W.; Wamwayi, H.M.; Hockley, E.; Shalla, M.S.; Goatley, L.; Knowles, N.J. J. Gen. Virol. 74, 275-2780, 1993 A;Reference number: P00865; MUID:94103786; PMID:8277286 A;Accession: P00865 A;Molecule type: mRNA A;Residues: 86-191 <CHA> A;Genetics: F C;Superfamily: parainfluenza virus cell fusion protein C;Keywords: glycoprotein; membrane fusion; transmembrane protein Query Match Score 72; DB 2; Length 546; Best Local Similarity 100.0%; Pred. No. 0.033; Mismatches 0; Indels 0; Gaps 0; Matches 15; Conservative 15; Query 3 LSEIKGVIVHLREGV 17 Db 284 LSEIKGVIVHLREGV 298

RESULT 5

B48556 cell fusion glycoprotein precursor - measles virus (strain AIK-C) C;Species: measles virus C;Accession: B48556 C;Date: 17-Feb-1994 #text_change 16-Jul-1999 R;Mori, T.; Sasaki, K.; Hashimoto, H.; Makino, S. Virus Genes 7, 67-81, 1993 A;Title: Molecular cloning and complete nucleotide sequence of genomic RNA of the AIK-C A;Reference number: B48556 A;Accession: B48556 A;Molecule type: Genomic RNA A;Residues: 1-550 <MOR> A;Cross-references: GB:S58435; NID:G299460; PIDN:JAB26145.1; PID:9299465 A;Note: sequence extracted from NCBI backbone (NCBIn:129264; NCBI:129272) C;Genetics: A;Gene: F C;Superfamily: parainfluenza virus cell fusion protein C;Keywords: glycoprotein; membrane fusion; transmembrane protein C;KeyWords: membrane fusion protein; transmembrane protein C;1-22/Domain: signal sequence #status predicted <SG> F;23-107/Product: cell fusion glycoprotein F2 #status predicted <FF2> F;108-550/Product: cell fusion glycoprotein F1 #status predicted <FF1> F;49-514/Domain: hydrophobic transmembrane #status predicted <TM> F;6,29,61,67/Binding site: carbohydrate (Asn) (covalent) #status predicted <FF1> Query Match Score 72; DB 1; Length 550; Best Local Similarity 100.0%; Pred. No. 0.033; Mismatches 0; Indels 0; Gaps 0; Matches 15; Conservative 15; Query 3 LSEIKGVIVHLREGV 17 Db 288 LSEIKGVIVHLREGV 302

RESULT 6

VGNZMV cell fusion glycoprotein precursor - measles virus C;Species: measles virus C;Accession: B26962; A25616; PQ0384 C;Date: 31-Mar-1988 #sequence_revision 31-Mar-1989 #text_change 16-Jun-2000 R;Buckland, R.; Gerald, C.; Barker, R.; Wild, T.F. J. Gen. Virol. 68, 1695-1703, 1987 A;Title: Fusion glycoprotein of measles virus: nucleotide sequence of the gene and complement A;Reference number: A92794; MUID:87224816; PMID:3585281 A;Accession: A26962 A;Molecule type: mRNA A;Residues: 1-53 <NUC> A;Cross-references: GB:D000090; NID:9222061; PIDN:BAA00056.1; PID:9222062 A;Experimental source: strain Halle A;Richardson, C.; Hull, D.; Greer, P.; Hasel, K.; Berkovich, A.; Englund, G.; Bellini, Virology 155, 508-523, 1986 A;Title: The nucleotide sequence of the mRNA encoding the fusion protein of measles virus A;Reference number: A94350; MUID:8707166; PMID:378802 A;Accession: A269616 A;Molecule type: mRNA A;Residues: 4-53 <RIC> A;Cross-references: GB:MI4915; NID:9331762; PIDN:AAA46423.1; PID:9331763 A;Experimental source: strain Edmonston R;Schulz, T.F.; Hood, J.G.; Whirby, D.; Tizard, E.J.; Dillon, M.J.; Weiss, R.A. J. Gen. Virol. 73, 1581-1586, 1992 A;Title: A measles virus isolate from a child with Kawasaki disease: sequence comparison A;Reference number: P00380 A;Molecule type: Genomic RNA A;Residues: 272-553 <SCH1> A;Experimental source: isolate CL A;Accession: P00384 A;Molecule type: Genomic RNA A;Residues: 272-553 <SCH2> A;Experimental source: isolate SE C;Genetics: F C;Superfamily: parainfluenza virus cell fusion protein

C;Keywords: glycoprotein; membrane fusion; transmembrane protein
 F;25/10/Domain: signal sequence #status predicted <SIG>
 F;111-553/Product: cell fusion glycoprotein F2 #status predicted <FF2>
 F;501-517/Domain: transmembrane#status predicted <TMN>
 F;32,64,70/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 45.0%; Score 72; DB 1; Length 553;
 Best Local Similarity 100.0%; Pred. No. 0.033; Gaps 0;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLLEGV 17
 Db 291 LSEIKGVIVHRLLEGV 305

RESULT 7
 VCNZRK
 Cell fusion glycoprotein Precursor - rinderpest virus (strain Kabete O)
 N/Contains: fusion glycoprotein F1; fusion glycoprotein F2
 C;Species: rinderpest virus
 C;Accession: A31051
 R;Hsu, D.; Yamanaka, M.; Miller, J.; Dale, B.; Grubman, M.; Yilma, T.
 Virology 166, 149-153, 1988
 A;Title: Cloning of the fusion gene of rinderpest virus: comparative sequence analysis
 A;Reference number: A31051; MUID:88322864; PMID:3413983
 A;Accession: A31051
 A;Molecule type: genomic RNA
 A;Residues: 1-546 <HSU>
 C;Genetics:
 A;Gene: P
 C;Superfamily: parainfluenza virus cell fusion protein
 C;Keywords: glycoprotein; membrane fusion; transmembrane protein
 F;1-19/Domain: signal sequence #status predicted <SIG>
 F;109-108/Product: cell fusion glycoprotein F2 #status predicted <FF1>
 F;109-134/Product: cell fusion glycoprotein F1 #status predicted <FF2>
 F;491-513/Domain: transmembrane#status predicted <TMN1>
 F;25,57,63,518/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 44.4%; Score 71; DB 1; Length 546;
 Best Local Similarity 93.3%; Pred. No. 0.046; Gaps 0;
 Matches 14; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLLEGV 17
 Db 284 LSEIKGVIVHRLLEGV 298

RESULT 8
 S47305
 Gene F Protein - rinderpest virus
 C;Species: rinderpest virus
 C;Date: 20-Oct-1994 #sequence_revision 08-Sep-1995 #text_change 20-Sep-1999
 A;Accession: S47305; S47301
 R;Baron, M.D.; Barrett, T.
 Submitted to the EMBL Data Library, March 1994
 A;Description: The sequence of the N and L genes of Rinderpest virus, and the 50 and 30
 A;Reference number: S47283
 A;Accession: S47305
 A;Molecule type: mRNA
 A;Residues: 1-546 <BAR>
 C;Superfamily: parainfluenza virus cell fusion protein
 C;Keywords: transmembrane protein

Query Match 44.4%; Score 71; DB 2; Length 546;
 Best Local Similarity 93.3%; Pred. No. 0.046; Gaps 0;
 Matches 14; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLLEGV 17
 Db 284 LSEIKGVIVHRLLESV 298

RESULT 9
 S47034
 Cell fusion protein precursor - porpoise morbillivirus
 N/Alternative names: F protein
 C;Species: porpoise morbillivirus
 C;Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 24-Nov-1999
 C;Accession: S47034
 R;Bolt, G.; Gottschalck, B.; Blixenkrone-Moeller, M.; Welsh, M.J.;
 submitted to the EMBL Data Library, July 1994
 A;Description: Nucleotide sequence comparisons of the F and M genes of cetacean morbilli
 A;Reference number: S47034
 A;Accession: S47034
 A;Molecule type: mRNA
 A;Residues: 1-552 <BOI>
 A;Cross-references: EMBL:X80757; PID:G520639; PIDN:CAA56731.1; PID:G520640
 A;Experimental source: isolate Ulster 88
 A;Note: the source is designated as Cetacean morbillivirus
 C;Superfamily: parainfluenza virus cell fusion protein
 F;1-25/Domain: signal sequence #status predicted <SIG>
 F;26-552/Product: fusion protein #status predicted <MAT>
 Query Match 41.6%; Score 66.5; DB 2; Length 552;
 Best Local Similarity 61.5%; Pred. No. 0.2; Mismatches 1; Indels 7; Gaps 1;
 Matches 16; Conservative 1; Mismatches 2; Indels 7; Gaps 1;

Qy 3 LSEIKGVIVHRLLEGVGPSLHWSYGL 28
 Db 290 LSEVKGVIVHRLLEAV-----SYNL 308

RESULT 10
 VGNZRL
 Cell fusion Glycoprotein precursor - rinderpest virus (strain L)
 N/Contains: fusion glycoprotein F1; fusion glycoprotein F2
 C;Species: rinderpest virus
 C;Date: 30-Sep-1999 #sequence_revision 30-Sep-1999 #text_change 16-Jul-1999
 C;Accession: A28921
 R;Tsukiyama, K.; Yoshihikawa, Y.; Yananouchi, K.
 Virology 164, 523-530, 1988
 A;Title: Fusion glycoprotein (F) of rinderpest virus: entire nucleotide sequence of th
 A;Reference number: A28921; MUID:88219541; PMID:3285575
 A;Accession: A28921
 A;Molecule type: mRNA
 A;Cross-references: GB:M20870; PIDN:AAA47399.1; PID:G333898
 C;Genetics:
 A;Gene: F
 C;Superfamily: parainfluenza virus cell fusion protein
 C;Keywords: glycoprotein; membrane fusion; transmembrane protein
 F;1-20/Domain: signal sequence #status predicted <SIG>
 F;20-104/Product: cell fusion glycoprotein F2 #status predicted <FG2>
 F;105-146/Product: cell fusion glycoprotein F1 #status predicted <FG1>
 F;109-133/Domain: transmembrane#status predicted <TMN1>
 F;185-513/Domain: transmembrane #status predicted <TMN2>
 F;25,57,63/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 41.2%; Score 66; DB 1; Length 546;
 Best Local Similarity 93.3%; Pred. No. 0.23; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLLEGV 17
 Db 284 LSEIKGVIVHRLLESV 298

RESULT 11
 JQ223
 Cell fusion protein F0 precursor - phocine distemper virus
 N/Contains: F1 and F2 chains
 C;Species: phocine distemper virus

C;Date: 14-Jul-1994 #sequence_revision 14-Jul-1994 #text_change 24-Nov-1999
 C;Accession: JQ2223
 C;Cross-references: GB:L07075
 A;Note: the authors transcribed the codon ATC for residue 4 as Leu
 C;Comment: This fusion protein F0 is cleaved into F1 and F2 chains.
 A;Gene: F
 C;Superfamily: Parainfluenza virus cell fusion protein
 C;Keywords: Glycoprotein; membrane fusion transmembrane protein
 F;1-15/Domain: signal sequence #status predicted <SG>
 F;16-99/Product: F2 chain #status predicted <F2C>
 F;105-1542/Product: F1 chain #status predicted <F1C>
 F;46-512/Domain: transmembrane #status predicted <TM>
 F;21,53,59,397/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 40.6%; Score 65; DB 2; Length 542;
 Best Local Similarity 86.7%; Pred. No. 0.32; Mismatches 1; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLEGV 17
 Db 280 LSEVKGVIVHRLEAV 294

RESULT 12
 VGN2D
 Cell fusion glycoprotein precursor - canine distemper virus
 N;Contains: Fusion protein F1; fusion protein F2
 C;Species: canine distemper virus
 C;Accession: JS0321
 C;Cross-references: GB:M21849; NID:9323241; PID:AA42878.1; PID:9323242
 A;Gene: F
 C;Superfamily: Parainfluenza virus cell fusion protein
 C;Keywords: Glycoprotein; membrane fusion transmembrane protein
 F;1-135/Domain: signal sequence #status predicted <SG>
 F;136-224/Product: cell fusion glycoprotein F2 #status predicted <F2P>
 F;606-629/Domain: transmembrane #status predicted <TM>
 F;62,141,173,179,517/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 40.6%; Score 65; DB 1; Length 662;
 Best Local Similarity 86.7%; Pred. No. 0.4; Mismatches 1; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLEGV 17
 Db 400 LSEVKGVIVHRLEAV 414

RESULT 13
 S21382
 Cell fusion protein - canine distemper virus
 C;Species: canine distemper virus
 C;Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 24-Nov-1999

C;Accession: S21382
 R;Wild, T.F.; Bernard, A.; Spehner, D.; Villeval, D.; Drillien, R.
 R;Wisser, I.Y.G.; van der Heijden, R.W.J.; van de Bildt, M.W.G.; Kenter, M.J.H.; Oervell
 J. Gen. Virol. 74, 1989-1984, 1993
 A;Description: Vaccination of mice against canine distemper virus induced encephalitis
 A;Reference number: S21382
 A;Accession: S21382
 A;Status: preliminary
 A;Molecule type: genomic RNA
 A;Residues: 1-662 <WIL>
 A;Cross-references: ENBL:X55509; NID:958853; PID:CAA46481.1; PMID:958854
 C;Superfamily: parainfluenza virus cell fusion protein
 Query Match 40.6%; Score 65; DB 2; Length 662;
 Best Local Similarity 86.7%; Pred. No. 0.4; Mismatches 1; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLEGV 17
 Db 400 LSEVKGVIVHRLEAV 414

RESULT 14
 VGN2D
 Cell fusion glycoprotein precursor - phocine distemper virus
 N;Contains: Fusion protein F1; fusion protein F2
 C;Species: phocine distemper virus
 C;Accession: JQ1368
 R;Koevamets, J.; Blixenkrone-Moeller, M.; Sharma, B.; Oervell, C.; Norrby, E.
 J. Gen. Virol. 72, 2959-2967, 1991
 A;Title: The nucleotide sequence and deduced amino acid composition of the haemagglutinir
 A;Reference number: JQ1368; MUID:92113538; PMID:1765768
 A;Accession: JQ1368
 A;Molecule type: genomic RNA
 A;Residues: 1-631 <SKOV>
 A;Genetics:
 C;Gene: F
 C;Superfamily: parainfluenza virus cell fusion protein
 C;Keywords: Glycoprotein; membrane fusion transmembrane protein
 F;1-188/Product: cell fusion glycoprotein F2 #status predicted <FP2>
 F;189-106/Domain: transmembrane #status predicted <TM1>
 F;189-193/Region: cleavage processing #status predicted <PP1>
 F;194-631/Product: cell fusion glycoprotein F1 #status predicted <FP1>
 F;194-212/Domain: transmembrane #status predicted <TM2>
 F;575-595/Domain: transmembrane #status predicted <TM3>
 F;110,142,148,486/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 40.0%; Score 64; DB 1; Length 631;
 Best Local Similarity 80.0%; Pred. No. 0.52; Mismatches 2; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLEGV 17
 Db 369 LSEVKGVIVHRLEAV 383

RESULT 15
 A48346
 Cell fusion glycoprotein precursor - phocine distemper virus (strain Ulster/88)
 N;Contains: Fusion protein F1; fusion protein F2
 C;Species: phocine distemper virus
 C;Accession: AA8346
 R;Curran, M.D.; Lu, Y.J.; Rima, B.K.
 Arch. Virol. 126, 159-167, 1992
 A;Title: The fusion protein gene of phocine distemper virus: nucleotide and deduced ami
 A;Reference number: A48346; MUID:9238837; PMID:1524494
 A;Accession: AA8346
 A;Molecule type: mRNA
 A;Residues: 1-831 <CUR>
 A;Note: sequence extracted from NCBI backbone (NCBIN:113098, NCBI:113099)
 C;Genetics:
 A;Gene: F

C;Superfamily: parainfluenza virus cell fusion protein
C;Keywords: Glycoprotein; membrane fusion; transmembrane protein
F;1-88/Product: cell fusion glycoprotein; P2 #status predicted <FP2>
F;89-106/Domain: transmembrane #status predicted <TM1>
F;194-631/Product: cell fusion glycoprotein F1 #status predicted <FP1>
F;194-219/Domain: transmembrane #status predicted <TM2>
F;575-595/Domain: transmembrane #status predicted <TM3>
F;110-142/Binding site: carbohydrate (Asn) (covalent) #status predicted
F;110-142/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 40.0%; Score 64; DB 1; Length 631;
Best Local Similarity 80.0%; Pred. No. 0.52;
Matches 12; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Qy 3 LSEIKGVTVHRLGV 17
Db 369 LSEVKGVTVHRLAV 383

Search completed: March 10, 2004, 09:16:45
Job time : 10.7086 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: March 10, 2004 08:58:53 ; Search time 5.66926 Seconds

(without alignments)
 284.724 Million cell updates/sec

Title: US-09-848-834A-9
 Perfect score: 160
 Sequence: 1 KLISEIKGVVHRLLEGVEGPSLHNSYGLRPX 31

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141661 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_42:*

Pred. No. is the number of results predicted by chance to have a a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match	Length	DB ID	Description
1	72	45.0	534	1	VGLF MEASY	P26032 measles vir
2	72	45.0	546	1	VGLF_RINDB	P41360 rinderpest
3	72	45.0	550	1	VGLF_MEASA	P235973 measles vir
4	72	45.0	550	1	VGLF_MEAS	P08310 measles vir
5	71	44.4	546	1	VGLF_RINDR	P41336 rinderpest
6	66	41.2	546	1	VGLF_RINDR	P10864 rinderpest
7	65	40.6	546	1	VGLF_RINDR	P12574 rinderpest
8	65	40.6	662	1	VGLF_CDVO	P12569 canine dist
9	64	40.0	529	1	VGLF_MEASI	P26031 measles vir
10	64	40.0	631	1	VGLF_PHODV	P288816 phocine dis
11	57.5	35.9	91	1	GONI_PIG	P499921 sus scrofa
12	57.5	35.9	92	1	GONI_HUMAN	P01148 homo sapien
13	57	35.6	90	1	GONI_MOUSE	P13562 mus musculus
14	57	35.6	92	1	GONI_RAT	P07430 rattus norv
15	54	33.8	67	1	GONI_MACMU	P55247 macaca mulu
16	53	33.1	508	1	VGLG_IRN	P07933 infectious
17	52.5	32.8	90	1	GONI_RANCA	Q90yf3 rana catesbe
18	52	32.5	61	1	GONI_SHEBAU	Q28588 ovis aries
19	52	32.5	63	1	GONI_MESAU	Q91913 mesocricetus
20	52	32.5	89	1	GONI_XENLA	P45636 xenopus lae
21	52	32.5	92	1	GONI_TUGGB	Q93335 tubailla glis
22	52	32.5	379	1	PURK_BACSU	P12045 bacillus su
23	50.5	31.6	92	1	GONI_CAVPO	Q54713 cavia porce
24	50.5	31.6	99	1	GONI_DICLLA	Q91a10 dicentrae arch
25	49	30.6	169	1	CX41_THUB	Q9i810 thunnus obe
26	49	30.6	582	1	SYD_CHLUTR	Q84546 chlamydia t
27	48.5	30.3	95	1	GONI_MORSA	Q73812 morone saxa
28	48	30.0	10	1	GONI_ALIMI	P37041 alligator m
29	48	30.0	92	1	GONI_CHICK	P37042 gallus galli
30	48	30.0	94	1	GONI_HABU	P51918 hapolochromi
31	48	30.0	95	1	GONI_PAGMA	P70074 pegas major
32	48	30.0	95	1	GONI_SPAAU	P51919 aparus aura
33	48	30.0	124	1	UCNI_HUMAN	P50509 homo sapien

ALIGNMENTS

34	48	30.0	576	1	DFA1_ANASP	Q8ynws anaebaena sp
35	47	29.4	512	1	UCNI_MOUSE	P81615 mus musculus
36	47	29.4	452	1	HEMN_RHOH	P33770 rhodobacter
37	47	29.4	452	1	DFA1_STNV3	P95651 rhodobacter
38	47	29.4	573	1	DFA2_ANASP	Q55393 synchocystis
39	47	29.4	579	1	TER2_ECOLI	Q8z0co anaebaena sp
40	46.5	29.1	207	1	RFL_PYRAB	P04483 escherichia
41	46.5	29.1	417	1	YHBJ_ACTAC	P9v151 pyrococcus
42	46	28.7	110	1	P96769 actinobacil	P12605 human parv
43	46	28.7	555	1	GSH1_P11HC	Q9nf6 onchocerca
44	46	28.7	652	1	DCUP_AQUAE	Q66667 aquifex aeo
45	45.5	28.4	338	1		

RESULT 1

ID_VGLF_MEASY	STANDARD;	PRT;	534 AA.
AC_P26032;			
DT_01-MAY-1992 (Rel. 22, Created)			
DT_01-MAY-1992 (Rel. 22, Last sequence update)			
DT_16-OCT-2001 (Rel. 40, Last annotation update)			
DE_Fusion glycoprotein precursor [Contains: Fusion glycoprotein F2; Fusion glycoprotein F1].			
DB_F.			
GN_Measles virus (strain Yamagata-1) (Subacute sclerose panencephalitis virus).			
OS_Viruses; ssRNA negative-strand viruses; Mononegavirales; Paramyxoviridae; Paramyxovirinae; Morbillivirus.			
OX_NCBI_TaxID=11239;			
RN_			
RP_SEQUENCE FROM N.A.			
RX_MEDLINE=9085702; PubMed=1698327;			
RA_Konase K., Haga T., Yoshikawa Y.; Sato T.A., Yamamotochi K.,			
RT_Molecular analysis of structural protein genes of the Yamagata-1 strain of defective subacute sclerosing Panencephalitis virus.			
RL_Virus Genes 4:173-181(1990).			
CC_-!_FUNCTION: This protein directs fusion of viral and cellular membranes			
CC_-!_SUBUNIT: THE MATURE FORM IS A DIMER OF POLYPEPTIDES F-1 AND F-2 LINKED BY A DISULFIDE BOND.			
CC_-!_SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein family.			
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CC_			
CC_BA01405.1; -			
DR_HSPB; P04847; 1SVE.			
DR_InterPro; IPR00776; Fusion_gly.			
DR_PFO0533; Fusion_protein.			
KW_Glycoprotein; Fusion_protein; Transmembrane; Envelope_protein; Signal			
CC_EMEL; D10546; BAA01405.1; -			
DR_HSPB; P04847; 1SVE.			
DR_InterPro; IPR00776; Fusion_gly.			
DR_PFO0533; Fusion_protein.			
KW_Glycoprotein; Fusion_protein; Transmembrane; Envelope_protein; Signal			
FT_SIGNAL	1	23	
FT_CHAIN	24	534	
FT_CHAIN	24	112	
FT_CHAIN	113	534	
FT_TRANSMEM	113	136	
FT_DOMAIN	137	494	
FT_DOMAIN	137	495	
FT_DOMAIN	515	534	
FT_DOMAIN	516	534	
FT_DOMAIN	68	195	
FT_DISULFID	29	29	
FT_DISULFID	61	61	
FT_CARBOHYD	67	67	
FT_CARBOHYD	534 AA;	57963 MW;	F5B21757E43844D CRC641;

DT 01-AUG-1988 (Rel. 08, Created)
 DT 01-AUG-1988 (Rel. 08, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Fusion glycoprotein precursor [Contains: Fusion glycoprotein F2; Fusion Glycoprotein F1].
 GN Measles virus (strain Edmonston) (Subacute sclerose panencephalitis virus, strain Halle) (Subacute sclerose panencephalitis virus, strain Leningrad-16) (Subacute sclerose panencephalitis virus, strain Philadelphia-26) (Subacute sclerose panencephalitis virus, strain Philadelphia-16) (Subacute sclerose panencephalitis virus).
 OS Measles virus (strain Edmonston-Zagreb) (Subacute sclerose panencephalitis virus, strain Philadelphia-26) (Subacute sclerose panencephalitis virus, and Measles virus (strain Edmonston B) (Subacute sclerose panencephalitis virus).
 OC Paramyxoviridae; Paramyxovirinae; Morbilliviruses.
 OC NCBI_TaxID=11235, 11236, 70147, 70148, 70149, 70146;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN Edmonston;
 RX MEDLINE=87071668, PubMed=3788062;
 RX MEDLINE=87224816, PubMed=1585281;
 RA Richardson C.D., Hull D., Greer P., Hasel K., Berkovich A., Englund G., Bellini W.J., Rima B., Lazzarini R.A.;
 RT "The nucleotide sequence of the mRNA encoding the fusion protein of measles virus (Edmonston strain): a comparison of fusion proteins from several different paramyxoviruses.",
 RT Virology 155:508-523 (1986).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN Halle;
 RX MEDLINE=87224816, PubMed=1585281;
 RA Buckland R., Gerald C., Barker R., Wild T.F.;
 RT "Fusion glycoprotein of measles virus: nucleotide sequence of the gene and comparison with other paramyxoviruses.",
 RL J. Gen. Virol. 66:1695-1703 (1987).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN Edmonston;
 RX MEDLINE=90085790, PubMed=2596022;
 RA Cattaneo R., Schmid A., Splechtauer P., Kaelin K., Bacsko K., Meulen V., Pardowitz J., Pranagan S., Rima B.K., Udem S.A.;
 RT "Mutated and hypermutated genes of persistent measles viruses which caused lethal human brain diseases.",
 RL Virology 173:415-425 (1989).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN Edmonston;
 RX MEDLINE=92263801, PubMed=1585658;
 RA Schmid A., Splechtauer P., Cattaneo R., Bacsko K., Ter Meulen V., Billerter M.A.;
 RT "Subacute sclerosing panencephalitis is typically characterized by alterations in the fusion protein cytoplasmic domain of the persisting measles virus.",
 RL Virology 188:910-915 (1992).
 RN [5]
 RP SEQUENCE FROM N.A.
 RC STRAIN Edmonston, Leningrad-16, and Edmonston-Zagreb;
 RX MEDLINE=94249283, PubMed=8191786;
 RA Rota J.S., Wang Z.D., Rota P.A., Bellini W.J.;
 RT "Comparison of sequences of the H, F, and N coding genes of measles virus vaccine strains.",
 RT Virus Res. 31:317-330 (1994).
 RN [6]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Philadelphia-26;
 RX MEDLINE=9430181, PubMed=8030232;
 RA Hummel K.B., Vanchiere J.A., Bellini W.J.;
 RT "Restriction of fusion protein mRNA as a mechanism of measles virus persistence.",

RL Virology 202:665-672 (1994).
 RN [7]
 RP SEQUENCE FROM N.A.
 RC STRAIN Edmonston B;
 RA Billerter M.A.;
 RL Submitted (Oct-1995) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: This protein directs fusion of viral and cellular membranes.
 CC -1- SUBUNIT: THE MATURE FORM IS A DIMER OF POLYPEPTIDES F-1 AND F-2 LINKED BY A DISULFIDE BOND.
 CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein family.
 CC
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 CC
 DR M14915 AAA46423 1 -
 DR EMBL; X05597; CAI29090_1; ALT_INIT.
 DR EMBL; X01711; AAA75498_1; ALT_INIT.
 DR EMBL; X01711; AAA75499_1; -
 DR EMBL; U03657; AAA56644_1; ALT_INIT.
 DR EMBL; U03657; AAA56649_1; ALT_INIT.
 DR EMBL; U03670; AAA56650_1; ALT_INIT.
 DR EMBL; U08416; AAA50550_1; ALT_INIT.
 DR EMBL; U266517; CAI91367_1; ALT_INIT.
 DR EMBL; U266517; CAI91368_1; -
 DR HSPP; P04849; 1SVP.
 DR InterPro; IP000716; Fusion_gly.
 DR Pfam; PF00523; fusion_gly_1.
 KW Glycoprotein; Fusion Protein; Transmembrane; Envelope protein; Signal.
 FT SIGNAL 1 23
 PT CHAIN 24 112
 PT CHAIN 24 112
 PT CHAIN 113 550
 PT TRANSMEM 113 550
 PT TRANSMEM 136 494
 PT DOMAIN 137 494
 PT DOMAIN 515 515
 PT DISULFID 516 550
 PT DISULFID 68 195
 PT CARBOHYD 29 29
 PT CARBOHYD 61 61
 PT CARBOHYD 67 67
 SQ SEQUENCE 550 AA; 59532 MW; TA44FICA82169093 CRC64;
 SQ
 Query Match 45.0% Score 72; DB 1; Length 550;
 Best Local Similarity 100.0%; Pred. No. 0.0062;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLIEGV 17
 Db 288 LSEIKGVIVHRLIEGV 302
 RESULT 5
 VGLF RINDR STANDARD PRT; 546 AA.
 ID VGLF RINDR STANDARD PRT; 546 AA.
 AC P4156;
 DT 01-FEB-1995 (Rel. 31, Created)
 DT 01-FEB-1995 (Rel. 31, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DB Fusion glycoprotein precursor [Contains: Fusion glycoprotein F2; Fusion glycoprotein F1].
 GN Rinderpest virus (strain RBOK) (RDV).
 OS Viruses; ssRNA negative-strand viruses; Mononegavirales;
 OC Paronyxoviridae; Paramyxovirinae; Morbilliviruses.
 OC NCBI_TaxID=336409;
 RN [1]

SEQUENCE FROM N.A.	PUBMED=7908860;	PubMed=7908860;
RX		
RA	Evans S.A., Baron M.D., Chamberlain R.W., Goatley L., Barrett T.;	
PT	"Nucleotide sequence comparisons of the fusion protein gene from rinderpest and attenuated strains of rinderpest virus."	
RL	J. Gen. Virol. 75:3611-3617(1994).	
CC	-I- FUNCTION: This protein directs fusion of viral and cellular membranes.	membranes
CC	-I- SUBUNIT: THE MATURE FORM IS A DIMER OF POLYPEPTIDES F-1 AND F-2 LINKED BY A DISULFIDE BOND.	THE MATURE FORM IS A DIMER OF POLYPEPTIDES F-1 AND F-2 LINKED BY A DISULFIDE BOND.
CC	-I- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein family.	Belongs to the paramyxoviruses fusion glycoprotein family.
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CC	DR EMBL; 230700; CAB83186.1; -;	DR EMBL; 230700; CAB83186.1; -;
CC	DR EMBL; 230697; CAB83181.1; -;	DR EMBL; 230697; CAB83181.1; -;
CC	DR PIR; SA1305; S4705.	DR PIR; SA1305; S4705.
CC	DR InterPro; IP00076; Fusion_gly.	DR InterPro; IP00076; Fusion_gly.
CC	DR Pfam; PF0053; fusion_gly; 1.	DR Pfam; PF0053; fusion_gly; 1.
CC	DR Glycoprotein; Fusion_Protein; Transmembrane; Envelope protein; Signal.	DR Glycoprotein; Fusion_Protein; Transmembrane; Envelope protein; Signal.
CC	FT SIGNAL 1 19	FT SIGNAL 1 19
CC	FT CHAIN 20 546	FT CHAIN 20 546
CC	FT CARBOHYD 108 F2 PROTEIN.	FT CARBOHYD 108 F2 PROTEIN.
CC	FT CHAIN 109 546	FT CHAIN 109 546
CC	FT DOMAIN 104 108	FT DOMAIN 104 108
CC	FT TRANSMEM 109 133	FT TRANSMEM 109 133
CC	FT TRANSMEM 109 133	FT TRANSMEM 109 133
CC	FT DOMAIN 484 513	FT DOMAIN 484 513
CC	FT TRANSMEM 514 517	FT TRANSMEM 514 517
CC	FT DISULFID 64 191	FT DISULFID 64 191
CC	FT CARBOHYD 25 25	FT CARBOHYD 25 25
CC	FT CARBOHYD 57 57	FT CARBOHYD 57 57
CC	FT CARBOHYD 63 63	FT CARBOHYD 63 63
CC	FT CARBOHYD 518 518	FT CARBOHYD 518 518
CC	SQ SEQUENCE 546 AA; 58705 MW;	SQ SEQUENCE 546 AA; 58911 MW;
CC		Query Match Score 66; DB 1; Length 546;
CC	Best Local Similarity 93.3%; Pred. No. 0.048;	Best Local Similarity 93.3%; Pred. No. 0.048;
CC	Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;	Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
CC		RESULT 7
CC	VGLF_RINDL STANDARD; PRT; 546 AA.	VGLF_RINDL STANDARD; PRT; 546 AA.
CC	AC P10864.	AC P10864.
CC	DT 01-JUL-1989 (Rel. 11, Created)	DT 01-JUL-1989 (Rel. 11, Last sequence update)
CC	DT 01-JUL-1989 (Rel. 11, Last annotation update)	DT 01-OCT-1989 (Rel. 12, Last annotation update)
CC	DE Fusion glycoprotein precursor [Contains: Fusion glycoprotein F2; Fusion glycoprotein F1].	DE Fusion glycoprotein precursor [Contains: Fusion glycoprotein F2; Fusion glycoprotein F1].
CC	GN F.	GN F.
CC	OS Rinderpest virus (strain Kabete 0) (RDV).	OS Rinderpest virus (strain Kabete 0) (RDV).
CC	OC Viruses; ssRNA negative-strand viruses; Mononegavirales;	OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
CC	OC Paramyxoviridae; Paramyxovirinae; Morbilliviridae.	OC Paramyxoviridae; Paramyxovirinae; Morbilliviridae.
CC	NCBI_TAXID=11242;	NCBI_TAXID=11242;
CC	RP SEQUENCE FROM N.A.	RP SEQUENCE FROM N.A.
CC	RA Hsu D., Yamakawa M., Miller J., Dale B., Grubman M., Yilma T.;	RA Hsu D., Yamakawa M., Miller J., Dale B., Grubman M., Yilma T.;
CC	RT "Cloning of the fusion gene of rinderpest virus: comparative sequence analysis with other morbilliviruses."	RT "Cloning of the fusion gene of rinderpest virus: comparative sequence analysis with other morbilliviruses."
CC	RL Virology 166:149-153 (1988).	RL Virology 166:149-153 (1988).
CC	CC -I- FUNCTION: This protein directs fusion of viral and cellular membranes.	CC -I- FUNCTION: This protein directs fusion of viral and cellular membranes.
CC	-I- SUBUNIT: THE MATURE FORM IS A DIMER OF POLYPEPTIDES F-1 AND F-2 LINKED BY A DISULFIDE BOND.	-I- SUBUNIT: THE MATURE FORM IS A DIMER OF POLYPEPTIDES F-1 AND F-2 LINKED BY A DISULFIDE BOND.
CC	-I- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein family.	-I- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein family.
CC	This SWISS-PROT entry is copyright. It is produced through a collaborative effort.	This SWISS-PROT entry is copyright. It is produced through a collaborative effort.

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CC EMBL; M21514; AAA44786.1; -
 DR PIR; A31051; VGNZRK.
 DR HSSEP; P04849; 1SVE.
 DR InterPro; IPR00776; Fusion_gly.
 DR Pfam; PF00523; fusion_gly; 1.
 KW Glycoprotein; Fusion_Protein; Transmembrane; Envelope protein; Signal.
 FT SIGNAL 1 19 FUSION GLYCOPROTEIN F0.
 FT CHAIN 20 546 F2 PROTEIN.
 FT CHAIN 20 108 F1 PROTEIN.
 FT DOMAIN 109 546 ARG-RICH (BASIC).
 FT DOMAIN 109 108 ARG-RICH (BASIC).
 FT TRANSMEM 109 133 POTENTIAL.
 FT TRANSMEM 109 513 ARG/LYS-RICH (BASIC).
 FT DOMAIN 514 513 N-LINKED (GLCNAC . . .) (POTENTIAL).
 FT CARBOYD 64 191 N-LINKED (GLCNAC . . .) (POTENTIAL).
 FT CARBOYD 25 25 N-LINKED (GLCNAC . . .) (POTENTIAL).
 FT CARBOYD 57 57 N-LINKED (GLCNAC . . .) (POTENTIAL).
 FT CARBOYD 63 63 N-LINKED (GLCNAC . . .) (POTENTIAL).
 FT CARBOYD 518 518 N-LINKED (GLCNAC . . .) (POTENTIAL).
 SQ SEQUENCE 546 AA; 58662 MW; 478D74DC18BCPCP CRC64;

Query Match 40.6%; Score 65; DB 1; Length 546;
 Best Local Similarity 86.7%; Pred. No. 0.088; DB 1; Length 662;
 Matches 13; Conservative 1; Missmatches 1; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHLEGV 17
 Db 284 LSEIKGVIVHLEGV 298

RESULT 8
 ID _VGLF_CDVO STANDARD; PRT; 662 AA.
 AC P12563; Q55991; 12. Created)
 DT 01-OCT-1989 (Rel. 12. Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Fusion Glycoprotein Precursor [Contains: Fusion glycoprotein F2;
 DE Fusion Glycoprotein F1].
 GN P. canin distemper virus (strain Onderstepoort) (CDV).
 OS Viruses; ssRNA negative-strand viruses; Mononegavirales;
 OC Paramyxoviridae; Paramyxovirinae; Morbilliviridae;
 RN [1] SEQUENCE FROM N.A.
 RP MEDLINE=88129050; PubMed=3433924;
 RX Barrett T.; Clarke D.K.; Evans S.A.; Rima B.K.; Villeval D.; Drillet R.; Wild T.F.; Bernard A.; Spehner D.; Billerer M.A.; "Vaccination of mice against canine distemper virus-induced encephalitis with vaccinia virus recombinants encoding measles or canine distemper virus antigens."; Vaccine 11:438-444 (1993).
 RT "The nucleotide sequence of the gene encoding the F protein of canine distemper virus: a comparison of the deduced amino acid sequence with other paramyxoviruses."; Virus Res. 8:373-386 (1987).
 RN [2] SEQUENCE FROM N.A.
 RP MEDLINE=9322769; PubMed=470428;
 RX Wild T.F.; Bernard A.; Spehner D.; Villeval D.; Drillet R.; Billerer M.A.; "Subacute sclerosing panencephalitis is typically characterized by alterations in the fusion protein cytoplasmic domain of the persisting measles virus"; Virology 188:910-911(1992).
 RT "FUNCTION: This protein directs fusion of viral and cellular membranes.";
 RT "SUBUNIT: THE MATURE FORM IS A DIMER OF POLYPEPTIDES F-1 AND F-2 LINKED BY A DISULFIDE BOND.";
 RT "SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein family.";
 RN [1] SEQUENCE FROM N.A.
 RP MEDLINE=92263801; PubMed=1585658;
 RX Schmid A.; Spiehler P.; Cattaneo R.; Bacsko K.; Ter Meulen V.; Billerer M.A.; "Subacute sclerosing panencephalitis is typically characterized by alterations in the fusion protein cytoplasmic domain of the persisting measles virus"; Virology 188:910-911(1992).
 RT "FUNCTION: This protein directs fusion of viral and cellular membranes.";
 RT "SUBUNIT: THE MATURE FORM IS A DIMER OF POLYPEPTIDES F-1 AND F-2 LINKED BY A DISULFIDE BOND.";
 RT "SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein family.";

SEQUENCE OF 24-33; PubMed=494667;
RX MEDLINE=7114303; PubMed=494667;
RA Baba Y.; Matsuo H.; Schally A.V.;
RT "Structure of the Porcine LH- and FSH-releasing hormone. II. Confirmation of the Proposed structure by conventional sequential biochemical Biophys. Res. Commun. 44:459-463 (1971)."
RT SEQUENCE FROM N.A.;
RL RN
RN [3]
RP SYNTHESIS OF GONADOLIBERIN.
RX MEDLINE=72065376; PubMed=942726;
RA Matsuo H.; Arimura A.; Nair R.M.G.; Schally A.V.;
RT "Synthesis of the porcine LH- and FSH-releasing hormone by the solid-phase method.";
RL RN
RP SYNTHESIS OF GONADOLIBERIN.
RX MEDLINE=72117544; PubMed=946275;
RA Baba Y.; Arimura A.; Schally A.V.;
RT "On the tryptophan residue in porcine LH and FSH-releasing hormone.";
RL RN
CC -!- FUNCTION: Stimulates the secretion of gonadotropins; it stimulates the secretion of both luteinizing and follicle-stimulating hormones.
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- SIMILARITY: Belongs to the GnRH family.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires license agreement (see <http://www.isb-sib.ch/announce/> or send an email to license@isb-sib.ch).
CC DR EMBL; L32854; AA031066 1; -
DR InterPro; IPR00012; GnRH.
DR Pfam; PF00446; GnRH; 1.
DR PRINTS; PR01541; GONADOLIBRNI.
KW Cleavage on pair of basic residues: Hormone; Amidation; Hypothalamus; Placenta; Signal; Pyrrolidone carboxylic acid.
FT SIGNAL 1 23 PROGONDOLIBERIN I.
FT CHAIN 24 91 GONADOLIBERIN I.
FT PEPTIDE 24 33 GNRH-ASSOCIATED PEPTIDE I.
FT PEPTIDE 34 91 APPEARS TO BE ESSENTIAL FOR BIOLOGICAL ACT_SITE 26 26 ACTIVITY.
FT MOD_RES 24 24 PYRROLIDONE CARBOXYLIC ACID.
FT MOD_RES 33 33 AMIDATION (G-34 PROVIDE AMIDE GROUP).
SQ SEQUENCE 91 AA; 10090 MW; 834044F32DDAA99 CRC64;

Query Match 35.9%; Score 57.5; DB 1; Length 91;
Best Local Similarity 50.0%; Pred. No. 0.14;
Matches 15; Conservative 4; Mismatches 8; Indels 3; Gaps 1;

Qy 1 KLLSEKIVVHLEGGPSLWWSYGLRP 30
Db 6 KLLA---GULLITLICVVGCSQHWSYGLRP 32

RESULT 12
ID GONI_HUMAN STANDARD; PRT; 92 AA.
AC P0114B;
DT 21-APR-1986 (Rel: 01, Created)
DT 01-APR-1988 (Rel: 07, Last sequence update)
AC P0114B;
DT 28-FEB-2003 (Rel: 41, Last annotation update)
DE Progondoliberin I precursor [Contains: Gonadoliberin I (LH-RH I) (luteinizing hormone-releasing hormone I) (Gonadotropin-releasing hormone I) (GnRH I) (Gonadorelin); GnRH-associated Peptide I].
DE GNRH1 OR GNRH OR LHRH.

OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Cetarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.;
RX MEDLINE=89166682; PubMed=267139;
RA Hayflick J.S.; Adelman J.P.; Seeburg P.H.;
RT "The complete nucleotide sequence of the human gonadotropin-releasing hormone gene.";
RT Nucleic Acids Res. 17:6403-6403 (1989).
RN [2]
RP SEQUENCE FROM N.A.;
RX MEDLINE=86054338; PubMed=2867548;
RA Adelman J.P.; Mason A.J.; Hayflick J.S.; Seeburg P.H.;
RT "Isolation of the gene and hypothalamic cDNA for the common precursor of Gonadotropin-releasing hormone and prolactin release-inhibiting factor in human and rat";
RT Proc. Natl. Acad. Sci. U.S.A. 83:179-183 (1986).
RN [3]
RP SEQUENCE FROM N.A.; AND VARIANT SER-16.
RX MEDLINE=85012739; PubMed=6090951;
RA Seeburg P.H.; Adelman J.P.;
RT "Characterization of cDNA for precursor of human luteinizing hormone releasing hormone.";
RT Nature 311:662-668 (1984).
RN [4]
RP SEQUENCE OF 24-33.
RX MEDLINE=83126573; PubMed=6760865;
RA Tan L.; Rousseau P.;
RT "The chemical identity of the immunoreactive LHRH-like peptide biosynthesized in the human placenta.";
RL Biochem. Biophys. Res. Commun. 109:1061-1071 (1982).
RN VARIANT SER-16.
RP MEDLINE=99318093; PubMed=10391209;
RA Cavigili M.; Altshuler D.; Ireland J.; Sklar P.; Ardlie K.; Patil N.; Shaw N.; Lane C.R.; Lim E.P.; Kalyanaraman N.; Nemesh J.; Ziaugra L.; Friedland L.; Rolfe A.; Warrington J.; Lipschutz R.; Daley G.Q.; Lander E.S.;
RT "Characterization of single-nucleotide polymorphisms in coding regions of human genes.";
RL Nat. Genet. 22:231-238 (1999).
RN [6]
RP ERBATUM.
RA Cavigili M.; Altshuler D.; Ireland J.; Sklar P.; Ardlie K.; Patil N.; Shaw N.; Lane C.R.; Lim E.P.; Kalyanaraman N.; Nemesh J.; Ziaugra L.; Friedland L.; Rolfe A.; Warrington J.; Lipschutz R.; Daley G.Q.; Lander E.S.;
RA "FUNCTION: Stimulates the secretion of gonadotropins; it stimulates the secretion of both luteinizing and follicle-stimulating hormones."
CC -!- SUBCELLULAR LOCATION: Secreted.
CC -!- FUNCTION: Stimulates the secretion of gonadotropins; it stimulates the secretion of both luteinizing and follicle-stimulating hormones.
CC -!- PHARMACEUTICAL: Available under the names Factrel (Ayerst Labs.), Lutrelese or Lutrelle (Ferring Pharmaceuticals) and Relisorm (Serono).
CC -!- SIMILARITY: Belongs to the GnRH family.
CC -!- PHARMACEUTICAL: Available through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See <http://www.isb-sib.ch/announce/> or send an email to license@isb-sib.ch).
CC -!- PHARMACEUTICAL: CAA55252; 1;
DR EMBL; X12578; AAA35916; 1;
DR EMBL; X15215; CAA3285; 1;
DR PIR; S05408; REHUG;
DR Genew; HGNC:1419; GNRH1;
DR MM; 152760; -

GO; GO:0005625; C:soluble fraction; TAS.

GO; GO:0005183; F:luteinizing hormone-releasing factor activity; TAS.

DR GO; GO:0007267; P:cell-cell signaling; TAS.

DR GO; GO:0007275; P:development; TAS.

DR GO; GO:0009285; F:negative regulation of cell proliferation; TAS.

DR InterPro; IPR00479; Gonadoliberin I.

DR GO; GO:00107165; F:signal transduction; TAS.

DR InterPro; IPR00212; GnrH.

DR InterPro; IPR00479; Gonadoliberin I.

DR PROSITE; PS000473; GNRH_1.

KW Cleavage on pair of basic residues; Hormone; Amidation; Hypothalamus;

KW Placenta; Pharmaceutical; Signal; Polymorphism;

KW Placental carboxylic acid.

FT SIGNAL 1 23 PROGONADOLIBERIN I.

FT CHAIN 24 92 PROGONADOLIBERIN I.

FT PEPTIDE 24 33 GNRH-ASSOCIATED PEPTIDE I.

FT ACT_SITE 37 92 APPEARS TO BE ESSENTIAL FOR BIOLOGICAL

ACTIVITY.

FT MOD_RES 24 24 PYRROLIDONE CARBOXYLIC ACID.

FT VARIANT 33 33 AMIDATION (G-34 PROVIDE AMIDE GROUP).

FT MOD_RES 33 16 W->S (in dbSNP:6185)

FT /FTId=V013943.

SO SEQUENCE 92 AA; 10380 MW; 30A72231B076FA79 CRC64;

Query Match Score 57.5; DB 1; Length 92;

Best Local Similarity 80.0%; Pred. No. 0.14;

Matches 12; Conservative 0; Mismatches 2; Indels 1; Gaps 1; Result 14

QY 17 VEG_PSLHNSYGLRP 30

DB 18 VEGCSSQQHNSYGLRP 32

GN GONI_MOUSE STANDARD; PRT; 90 AA.

ID GONI_MOUSE PRT; 90 AA.

AC P13162; PRT; 90 AA.

DT 01-JAN-1990 (Rel. 13, Created)

DT 01-JAN-1990 (Rel. 13, Last sequence update)

DR Progonadoliberin I precursor [Contains: Gonadoliberin I (LH-RH I)]

DE (Luteinizing hormone-releasing hormone I) (Gonadotropin-releasing factor II) (GnRH I) (Liliberin I); Prolactin release-inhibiting factor

DE GNRH1 OR GNRH

GN Rattus norvegicus (Rat)

OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

OC NCBI_TaxID=10116;

RN [1]

RP SEQUENCE FROM N.A. PMID=8609438; PubMed=2867548;

RP SEQUENCE FROM N.A. PMID=8334861; PubMed=2476669;

RP Maier C.C., Marchetti B., Leboeuf R.D., Blalock J.E./

RT Thymocytes express a mRNA that is identical to hypothalamic

RT luteinizing hormone-releasing hormone mRNA.;

RT Cell. Mol. Neurobiol. 12:447-454 (1992).

RN [4]

RP SEQUENCE OF 1-47 FROM N.A.

RC TISSUE=Heart; PMID=87149087; PubMed=3547652;

RC Adelman J.P., Bond C.T., Douglass J., Herbert E.,

CC !- SUBCELLULAR LOCATION: Secreted.

CC !- SIMILARITY: Belongs to the GnRH family.

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"Two mammalian genes transcribed from opposite strands of the same DNA locus";
 RT Factor alpha and its receptor in the hypothalamus of female rhesus
 RL Neuroendocrinology 60:346-359(1994).

-!- FUNCTION: stimulates the secretion of gonadotropins; it stimulates the secretion of both luteinizing and follicle-stimulating hormones.

-!- SUBCELLULAR LOCATION: Secreted.

-!- TISSUE SPECIFICITY: Central nervous system.

-!- SIMILARITY: Belongs to the Gnrh family.

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CC or send an email to license@isb-sib.ch.

CC DR EMBL; S50870; ABB24572; 1;

CC DR EMBL; M12579; AAA41263; 1;

CC DR EMBL; M31670; AAA41264; 1;

CC DR EMBL; M15527; AAA42141; 1; ALT_SEQ.

CC DR EMBL; M15528; AAA42239; 1;

CC DR EMBL; M15529; -; NOT_ANNOTATED_CDS.

CC DR PIR; M40147; RHTG.

CC DR InterPro; IPR000212; Gnrh.

CC DR InterPro; IPR004079; Gonadoliberini.

CC DR PRINTS; PF00446; Gnrh; 1.

CC DR PRINTS; PRO1541; GONADOLIBERNI.

CC DR PROSITE; PS00471; Gnrh; 1.

CC KW Cleavage on Pair of basic residues; Hormone; Amidation; Hypothalamus;

CC KW Placenta; Signal; Pyrrolidone carboxylic acid.

CC FT SIGNAL 1 23 PROGONADOLIBERIN I.

CC FT CHAIN 24 92 GONADOLIBERIN I.

CC FT PEPTIDE 37 92 PROLACTIN RELEASE-INHIBITING FACTOR I.

CC FT ACT_SITE 26 26 APPEARS TO BE ESSENTIAL FOR BIOLOGICAL ACTIVITY.

CC FT MOD_RES 24 24 PYRROLIDONE CARBOXYLIC ACID.

CC FT MOD_RES 33 33 AMIDATION (G-34 PROVIDE AMIDE GROUP).

CC SQ SEQUENCE 92 AA; 10500 MW; 494B564D8A3B3 CRC64;

CC Query Match 35.6%; Score 57; DB 1; Length 92;

CC Best Local Similarity 70.6%; Pred. No. 0.17; 3; Indels 2; Gaps 1;

CC Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CC Qy 14 LEGVEGDSLHWSYGLRP 30

CC Db 18 LECSCS-SQHWSYGLRP 32

RESULT 15
 GON1_MACMU
 ID GON1_MACMU STANDARD; PRT; 67 AA.
 AC P55247;
 DT 01-OCT-1996 (Rel. 34; Created)
 DT 01-OCT-1996 (Rel. 34; Last sequence update)
 DT 28-FEB-2003 (Rel. 41; Last annotation update)

DE Progonadoliberin I precursor (Contains: Gonadoliberin I (LH-RH I) (luteinizing hormone-releasing hormone I) (Gonadotropin-releasing hormone I) (Gnrh I) (Luliberin I); Gnrh-associated peptide I (Fragment)).
 DE GNRH1 OR Gnrh OR LRRH.
 OS Macaca mulatta (Rhesus macaque).
 OC Mammalia; Eutheria; Primates; Catarrhini; Euteleostomi; Cercopithecoidea; Cercopithecinae; Macaca.
 OC NCBI_TAXID=9544;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE-Hypothalamus;
 RX MEDLINE=95114501; PubMed=7545971;
 RX Ma Y.J., Costa M.E., Ojeda S.R.,

Search completed: March 10, 2004, 09:13:52
 Job time : 5.66926 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using SW model

Run on: March 10, 2004, 08:58:54 ; Search time 30.6381 Seconds
Perfect score: 160 (without alignments)

319.245 Million cell updates/sec

Title: US-09-848-834a-9
Scoring table: BLASTM62
GapP 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Lassing first 45 summaries

Database : SPREMBL 25:
1: sp_archea:
2: sp_bacteria:
3: sp_fungi:
4: sp_human:
5: sp_invertebrate:
6: sp_mammal:
7: sp_mmc:
8: sp_organelle:
9: sp_phage:
10: sp_plant:
11: sp_rabbit:
12: sp_virus:
13: sp_vertebrate:
14: sp_unclassified:
15: sp_tvirus:
16: sp_bacteriophage:
17: sp_archaea:
18: sp_bacteria:
19: sp_fungi:
20: sp_invertebrate:
21: sp_mammal:
22: sp_rabbit:
23: sp_tvirus:
24: sp_vertebrate:
25: sp_unclassified:
26: sp_tvirus:
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371: sp_fungi:
372: sp_invertebrate:
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467: sp_bacteriophage:
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1021: sp_invertebrate:
1022: sp_mammal:
1023: sp_rabbit:
1024:

DR GO; GO:0006948; P:viral-induced cell-cell fusion; IEA.

DR InterPro; IPR00776; Fusion_gly.

DR Pfam; PF00523; fusion_gly_1; 0234C2BAE193E77D CRC64;

SEQUENCE 545 AA; 58307 MW;

RESULT 2

Q04242 PRELIMINARY; PRT; 537 AA.

ID Q04242; AC 01-NOV-1996 (TREMBLrel. 01, Created)

DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)

DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)

DE Fusion protein.

CN

OS Measles virus.

Virus; ssRNA negative-strand viruses; Mononegavirales;

OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.

NCB_TaxID=11234;

RN [1]

SEQUENCE FROM N.A.

RP SEQUENCE FROM N.A.

EMBL; S9003063; PubMed=31167982;

RA Cattaneo R., Schmid A., Eischle D., Bacsko K., ter Meulen V.,

RA Billeret M.A.;

RT "Biased hypermutation and other genetic changes in defective measles viruses in human brain infections.";

RT Cell 55:255-265(1988).

RN [2]

RP SEQUENCE FROM N.A.

RA Cattaneo R., Billeret M.A.;

RL Virology 0.; 0(0);

DR EMBL; X16567; CAA34574.1; -;

DR EMBL; X16567; CAA34575.1; -;

DR HSSP; P04849; ISVF.

GO; GO:0019039; P:viral-induced fusion molecule activity; IEA.

DR GO; GO:0006948; P:viral-induced cell-cell fusion; IEA.

DR InterPro; IPR00776; Fusion_gly.

DR Pfam; PF00523; fusion_gly_1;

SEQUENCE 537 AA; 58275 MW; DOA60AC66D97E06 CRC64;

Query Match

Best Local Similarity 100.0%; Score 72; DB 12; Length 537;

Matches 15; Conservative 0; Missmatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLEGV 17

Db 291 LSEIKGVIVHRLEGV 305

RESULT 3

Q9PXA4 PRELIMINARY; PRT; 545 AA.

ID Q9PXA4; AC 01-MAY-2000 (TREMBLrel. 13, Created)

DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)

DE Fusion protein.

OS Measles virus.

Virus; ssRNA negative-strand viruses; Mononegavirales;

OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.

NCB_TaxID=11234;

RN

RP SEQUENCE FROM N.A.

RC STRAIN=OSA-3;

RA Ning X., Ayata M., Morimoto K., Ito N., Shingai M., Kimura M., Ogura H.;

RT Nucleotide sequences of the fusion protein gene of subacute sclerosing panencephalitis viruses: deduced amino acid sequences showed the cytoplasmic domain highly mutated --truncated, elongated or predicted secondary structure changed.";

RT Submitted (AUG-1999) to the EMBL/GenBank/DDBJ databases.

RL EMBL; AF179440; AAF02704.1; -;

DR HSSP; P04849; ISVF.

DR GO; GO:0019039; P:viral-cell fusion molecule activity; IEA.

DR GO; GO:0006948; P:viral-induced cell-cell fusion; IEA.

DR InterPro; IPR00776; Fusion_gly_1;

DR Pfam; PF00523; fusion_gly_1;

SEQUENCE 546 AA; 58572 MW; 449B2B2DD7405F0B CRC64;

Query Match

Best Local Similarity 100.0%; Score 72; DB 12; Length 546;

Matches 15; Conservative 0; Missmatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLEGV 17

Db 284 LSEIKGVIVHRLEGV 298

RESULT 4

Q91HHS PRELIMINARY; PRT; 546 AA.

ID Q91HHS; AC 01-OCT-2001 (TREMBLrel. 19, Created)

DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)

DE Fusion protein.

GN F.

OS Rinderpest virus.

Virus; ssRNA negative-strand viruses; Mononegavirales;

OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.

NCB_TaxID=11241;

RN

RP SEQUENCE FROM N.A.

RC STRAIN=N.K.;

RX MEDLINE=21014265; PubMed=11186456;

RA Aianot P.K., Sminev A.G., Bezborodova S.V., Starov S.K., Drygin V.V., Gusev A.A.;

RT "Primary structure of the F-gene from Rinderpest virus strain K.," Mol. Gen. Mikrobiol. Virusol. 4:29-33 (2000).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=N.K.;

RA Aianot P.K., Sminev A.G., Bezborodova S.V., Starov S.K., Drygin V.V., Gusev A.A.;

RT Submitted (MAY-2001) to the EMBL/GenBank/DDBJ databases.

RL EMBL; AY035887; AAK63190.1; -.

DR PIR; P00865; P00866.

DR PIR; P00867; P00867.

DR PIR; P00873; P00873.

DR GO; GO:0019039; P:viral-cell fusion molecule activity; IEA.

DR GO; GO:0006948; P:viral-induced cell-cell fusion; IEA.

DR InterPro; IPR00776; Fusion_gly_1;

DR Pfam; PF00523; fusion_gly_1;

SEQUENCE 546 AA; 58572 MW;

Query Match

Best Local Similarity 100.0%; Score 72; DB 12; Length 546;

Matches 15; Conservative 0; Missmatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLEGV 17

Db 284 LSEIKGVIVHRLEGV 298

RESULT 5

P90331 PRELIMINARY; PRT; 550 AA.

ID P90331; AC 01-MAY-1997 (TREMBLrel. 03, Created)

DT 01-MAY-1999 (TREMBLrel. 03, Last sequence update)

DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)

DE Fusion protein.

GN F.

OS Measles virus.

Virus; ssRNA negative-strand viruses; Mononegavirales;

OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.

NCB_TaxID=11234;

[1]	SEQUENCE FROM N.A. STRAIN=NAGAHATA;	Query Match 45.0%; Score 72; DB 12; Length 550; Best Local Similarity 100.0%; Pred. No. 0.075; Mismatches 0; Indels 0; Gaps 0
RA RT RN	"Sheng J., Watanabe M., Ueda S.; Submitted (AUG-1995) to the EMBL/GenBank/DDBJ databases. [2]	RP SEQUENCE FROM N.A. STRAIN=NAGAHATA; Sheng J., Nakaniishi M., Watanabe M., Ueda S.; "An amino acid alteration of F protein responsible for the enhanced fusogenicity of measles virus."; Submitted (AUG-1995) to the EMBL/GenBank/DDBJ databases.
RA RT RN	"Ning X., Ayata M., Morimoto K., Ito N., Shingai M., Kimura M., Ogura H.; Ogura H.; "Nucleotide sequences of the fusion protein gene of subacute sclerosing panencephalitis viruses: deduced amino acid sequences showed the cytoplasmic domain highly mutated -truncated, elongated or predicted secondary structure changed."; Submitted (AUG-1999) to the EMBL/GenBank/DDBJ databases. EMBL; D61926; BA009958.1; -. DR DR PTR; AF179431; AF02696.1; -. DR DR HSSP; P00376; P00376. DR DR GO; GO:001939; ISVF. DR DR GO; GO:0006348; P: viral-induced cell-cell fusion; IEA. InterPro; IPR000776; Fusion_gly. Pfam; PF000523; AA; 59530 MW; 97C991C7E2169839 CR64; SQ SEQUENCE 550 AA;	RP SEQUENCE FROM N.A. STRAIN=NAGAHATA; Ning X., Ayata M., Morimoto K., Ito N., Shingai M., Kimura M., Ogura H.; "Nucleotide sequences of the fusion protein gene of subacute sclerosing panencephalitis viruses: deduced amino acid sequences showed the cytoplasmic domain highly mutated -truncated, elongated or predicted secondary structure changed."; Submitted (AUG-1999) to the EMBL/GenBank/DDBJ databases. EMBL; D61926; BA009958.1; -. DR DR PTR; AF179431; AF02696.1; -. DR DR HSSP; P00376; P00376. DR DR GO; GO:001939; ISVF. DR DR GO; GO:0006348; P: viral-induced cell-cell fusion; IEA. InterPro; IPR000776; Fusion_gly. Pfam; PF000523; AA; 59530 MW; 97C991C7E2169839 CR64; SQ SEQUENCE 550 AA;
RA RT RN	Query Match 45.0%; Score 72; DB 12; Length 550; Best Local Similarity 100.0%; Pred. No. 0.075; Mismatches 0; Indels 0; Gaps 0;	Query Match 45.0%; Score 72; DB 12; Length 550; Best Local Similarity 100.0%; Pred. No. 0.075; Mismatches 0; Indels 0; Gaps 0;
RA RT RN	Query Match 45.0%; Score 72; DB 12; Length 550; Best Local Similarity 100.0%; Pred. No. 0.075; Mismatches 0; Indels 0; Gaps 0;	Query Match 45.0%; Score 72; DB 12; Length 550; Best Local Similarity 100.0%; Pred. No. 0.075; Mismatches 0; Indels 0; Gaps 0;
RA RT RN	RESULT 6 Q9QEX0 PRELIMINARY; PRT; 550 AA. ID Q9QEX0; AC Q9QEX0; DT 01-MAY-2000 (TREMBLrel. 13. Created) DT 01-MAY-2000 (TREMBLrel. 13. Last sequence update) DT 01-OCT-2003 (TREMBLrel. 25. Last annotation update) DE Fusion protein. OS Measles virus. Viruses; ssRNA, negative-strand viruses; Mononegavirales; Paramyxoviridae; Paramyxovirinae; Morbillivirus. [1] NCBI_TaxID=11234; RN RP SEQUENCE FROM N.A. STRAIN=Nagahata (N.A.); Ning X., Ayata M., Morimoto K., Ito N., Shingai M., Kimura M., Ogura H.; "Nucleotide sequences of the fusion protein gene of subacute sclerosing panencephalitis viruses: deduced amino acid sequences showed the cytoplasmic domain highly mutated -truncated, elongated or predicted secondary structure changed."; Submitted (AUG-1999) to the EMBL/GenBank/DDBJ databases. HSSP; P00376; P00376. Pfam; AF179432; AF02697.1; -. DR DR GO; GO:0019039; ISVF. DR DR GO; GO:0006348; P: viral-induced cell-cell fusion; IEA. InterPro; IPR000776; Fusion_gly. Pfam; PF000523; AA; 59530 MW; 2AA969D37FA5CA17 CRC64; SQ SEQUENCE 550 AA;	RESULT 7 Q9QE99 PRELIMINARY; PRT; 550 AA. ID Q9QE99; AC Q9QE99; DT 01-MAY-2000 (TREMBLrel. 13. Created) DT 01-MAY-2000 (TREMBLrel. 13. Last sequence update) DT 01-OCT-2003 (TREMBLrel. 25. Last annotation update) DE Fusion protein. OS Measles virus. Viruses; ssRNA, negative-strand viruses; Mononegavirales; Paramyxoviridae; Paramyxovirinae; Morbillivirus. [1] NCBI_TaxID=11234; RN RP SEQUENCE FROM N.A. STRAIN=Nagahata (N.A.); Ning X., Ayata M., Morimoto K., Ito N., Shingai M., Kimura M., Ogura H.; "Nucleotide sequences of the fusion protein gene of subacute sclerosing panencephalitis viruses: deduced amino acid sequences showed the cytoplasmic domain highly mutated -truncated, elongated or predicted secondary structure changed."; Submitted (AUG-1999) to the EMBL/GenBank/DDBJ databases. HSSP; P00376; P00376. Pfam; AF179432; AF02697.1; -. DR DR GO; GO:0019039; ISVF. DR DR GO; GO:0006348; P: viral-induced cell-cell fusion; IEA. InterPro; IPR000776; Fusion_gly. Pfam; PF000523; AA; 59530 MW; 2AA969D37FA5CA17 CRC64;

RT fusogenicity of measles virus."; to the EMBL/GenBank/NCBI databases.

RL Submitted (AUG-1995) to the EMBL/GenBank/NCBI databases.

DR EMBL: D63924; BRA09951.1; -.

DR PIR: PQ0376; PQ0316.

DR HSSP: P04849; ISVF.

DR GO; GO:0019039; F:viral-cell fusion molecule activity; IEA.

DR GO; GO:0006848; F:viral-induced cell-cell fusion; IEA.

DR InterPro: IPR000776; Fusion_gly.

DR Pfam: PF00523; Fusion_gly; T.

SEQUENCE 550 AA; 5589 MW; 73E7BD457ABA39B7 CRC64;

Query Match 45.0%; Score 72; DB 12; Length 550;

Best Local Similarity 100.0%; Pred. No. 0.075; Mismatches 0; Gaps 0; Indels 0; Gaps 0;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLEGV 17

Db 288 LSEIKGVIVHRLEGV 302

RESULT 9

Q9QW77 PRELIMINARY; PRT; 550 AA.

AC Q9QW77; PRELIMINARY; PRT; 550 AA.

DT 01-MAY-2000 (TREMBLrel. 13, Created)

DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)

DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)

DE Fusion protein.

OS Measles virus.

OC Viruses; ssRNA negative-strand viruses; Mononegavirales;

OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.

NCBI_TaxID=11234;

RN SEQUENCE FROM N.A.

STRAIN=OSA-2;

RA Ning X., Ayata M., Morimoto K., Ito N., Shingai M., Kimura M., Ogura H.,

RT "Nucleotide sequences of the fusion protein gene of subacute sclerosing panencephalitis viruses: deduced amino acid sequences showed the cytoplasmic domain highly mutated - truncated or predicted secondary structure changed."; to the EMBL/GenBank/NCBI databases.

DR EMBL: AF179435; AAC02703.1; -.

DR PIR: PQ0376; PQ0376.

DR HSSP; P04849; ISVF.

DR GO; GO:0019039; F:viral-cell fusion molecule activity; IEA.

DR GO; GO:0006848; F:viral-induced cell-cell fusion; IEA.

DR InterPro: IPR000776; Fusion_gly.

DR Pfam: PF00523; Fusion_gly; T.

SEQUENCE 550 AA; 5533 MW; 086E51FED5582BA CRC64;

Query Match 45.0%; Score 72; DB 12; Length 550;

Best Local Similarity 100.0%; Pred. No. 0.075; Mismatches 0; Gaps 0; Indels 0; Gaps 0;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DE Fusion protein.

GN P.

Q9WK44 PRELIMINARY; PRT; 550 AA.

AC Q9WK44; PRELIMINARY; PRT; 550 AA.

DT 01-NOV-1999 (TREMBLrel. 12, Created)

DT 01-NOV-1999 (TREMBLrel. 12, Last sequence update)

DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)

DE Fusion protein.

OS Measles virus.

OC Viruses; ssRNA negative-strand viruses; Mononegavirales;

OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.

NCBI_TaxID=11234;

RN [1] SEQUENCE FROM N.A.

RP STRAIN=MTC; PubMed=10400788;

RC MEDLINE=99329215; PubMed=10400788;

RX Johnstone I.C.; Ter Meulen V.; Schneider-Schaubies J., Schneider-Schaubies S.; vaccine virus expressing wild-type glycoprotein; consequences for viral spread and cell tropism. J. Virol. 73:6903-6915(1999).

RA RT "A recombinant measles vaccine virus expressing wild-type glycoprotein: consequences for viral spread and cell tropism."; J. Virol. 73:6903-6915(1999).

RA RL AJ133108; CAB38075.1; -.

RA DR P00376; PQ0376.

RA DR HSSP; P04849; ISVF.

RA DR GO:0006948; F:viral-cell fusion molecule activity; IEA.

RA DR GO:0019039; F:viral-induced cell-cell fusion; IEA.

RA DR InterPro: IPR000776; Fusion_gly.

RA DR Pfam: PF00523; Fusion_gly; T.

RA SQ SEQUENCE 550 AA; 59580 MW; 8255499968B5D862 CRC64;

Query Match 45.0%; Score 72; DB 12; Length 550;

Best Local Similarity 100.0%; Pred. No. 0.075; Mismatches 0; Indels 0; Gaps 0;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLEGV 17

Db 288 LSEIKGVIVHRLEGV 302

RESULT 11

Q89495 PRELIMINARY; PRT; 550 AA.

AC Q89495; PRELIMINARY; PRT; 550 AA.

DT 01-NOV-1996 (TREMBLrel. 01, Created)

DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)

DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)

DE Fusion protein.

GN P.

OS Measles virus.

OC Viruses; ssRNA negative-strand viruses; Mononegavirales;

OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.

NCBI_TaxID=11234;

RN SEQUENCE FROM N.A.

STRAIN=OSA-2;

RA Rota J.S., Hummel K.B., Rota P.A., Bellini W.J.; "Genetic variability of the glycoprotein genes of current wild-type measles isolates"; J. Virol. 188:35-42(1992).

RA Rota J.S., Hummel K.B., Rota P.A., Bellini W.J.; "Genetic variability of the glycoprotein genes of current wild-type measles isolates"; J. Virol. 188:35-42(1992).

RA DR EMBL: MB1903; AAA46421.1; -.

RA DR EMBL: MB1901; AAA46421.1; -.

RA DR PIR: PQ0376; PQ0376.

RA DR HSSP; P04849; ISVF.

RA DR GO:0019039; F:viral-cell fusion molecule activity; IEA.

RA DR GO:0006948; F:viral-induced cell-cell fusion; IEA.

RA DR InterPro: IPR000776; Fusion_gly.

RA DR Pfam: PF00523; Fusion_gly; T.

RA SQ SEQUENCE 550 AA; 53364 MW; A78EEC9CD6268E58 CRC64;

Query Match 45.0%; Score 72; DB 12; Length 550;

Best Local Similarity 100.0%; Pred. No. 0.075; Mismatches 0; Indels 0; Gaps 0;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLEGV 17

Db 288 LSEIKGVIVHRLEGV 302

RESULT 12

Q8V049 PRELIMINARY; PRT; 550 AA.

AC Q8V049; PRELIMINARY; PRT; 550 AA.

DT 01-MAR-2002 (TREMBLrel. 20, Created)

DT 01-MAR-2002 (TREMBLrel. 20, Last sequence update)

DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)

DE Fusion protein.
 F. Measles virus.
 OS Mononegavirales; Morbillivirus.
 OC ssRNA negative-strand viruses; Paramyxovirinae; Morbillivirus.
 OC Paramyxoviridae; Paramyxovirinae;
 OC NCBI_TaxID=11234;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=G954;
 RX MEDLINE=21635526; PubMed=11773423;
 RA Waku Kouenou D., Wild T.F.;
 RT "Adaptation of wild-type measles virus to tissue culture.";
 RL EMBL:AY059392; AAL29688.1; -.
 DR PIR; PQ0376; PQ0376.
 DR GO; GO:001039; P:viral-cell fusion molecule activity; IEA.
 DR GO; GO:0026948; P:viral-induced cell-cell fusion; IEA.
 DR InterPro; IPR000776; Fusion_gly.
 DR Pfam; PF00523; fusion_gly; MW; 9A7A4BA9E4DA8E9 CRC64;
 SQ SEQUENCE 550 AA; 59551 MW;

Query Match 45.0%; Score 72; DB 12; Length 550;
 Best Local Similarity 100.0%; Pred. No. 0.075; Indels 0; Gaps 0;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLLEGV 17
 Db 288 LSEIKGVIVHRLLEGV 302

RESULT 13
 C9YJ94 PRELIMINARY; PRT; 550 AA.
 ID Q9YJ94;
 AC Q9YJ94;
 DT 01-MAY-1999 (TREMBLrel. 10, Created)
 DT 01-MAY-1999 (TREMBLrel. 10, Last sequence update)
 DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
 DE Fusion protein.
 OS Measles virus.
 OC ssRNA negative-strand viruses; Mononegavirales;
 OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.
 OC NCBI_TaxID=11234;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=9301V;
 RX MEDLINE=98440528; PubMed=9765110;
 RA Takeda M., Kato A., Kobune F., Sakata H., Li Y., Shioda T., Sakai Y., Asakawa M., Nagai Y.;
 RT "Measles virus attenuation associated with transcriptional impediment and a few amino acid changes in the polymerase and accessory proteins."
 RL J Virol. 72: 8650-8656 (1998).
 DR EMBL; AB012949; BAA3877.1; -.
 DR EMBL; AB012948; BAA3877.1; -.
 DR PIR; PQ0376; PQ0376.
 DR HSSP; PQ04849; 1SVP.
 DR GO; GO:0019039; P:viral-cell fusion molecule activity; IEA.
 DR InterPro; IPR000776; Fusion_gly.
 DR Pfam; PF00523; fusion_gly; MW; 7AA4FD117197BF9 CRC64;
 SQ SEQUENCE 550 AA; 59512 MW;

Query Match 45.0%; Score 72; DB 12; Length 550;
 Best Local Similarity 100.0%; Pred. No. 0.075; Indels 0; Gaps 0;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLLEGV 17
 Db 288 LSEIKGVIVHRLLEGV 302

RESULT 14
 C9QEX1 PRELIMINARY; PRT; 550 AA.
 ID Q9QEX1;
 AC Q9QEX1;
 DT 01-MAY-2000 (TREMBLrel. 13, Created)
 DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
 DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
 DE Fusion protein.
 OS Measles virus.
 OC ssRNA negative-strand viruses; Mononegavirales;
 OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.
 OC NCBI_TaxID=11234;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Matsuoka;
 RA Ning X., Ayata M., Morimoto K., Ito N., Shingai M., Kimura M., Ogura H.;
 RT "Nucleotide sequences of the fusion protein gene of subacute sclerosing panencephalitis viruses: deduced amino acid sequences showed the cytoplasmic domain highly mutated --truncated, elongated or predicted secondary structure changed.";
 RT Submitted (AUG-1999) to the EMBL/GenBank/DDBJ databases.
 RL EMBL; AF179430; AAF026951; -.
 DR PIR; PQ0376; PQ0376.
 DR HSSP; P04849; 1SVP.
 DR GO; GO:0019039; P:viral-cell fusion molecule activity; IEA.
 DR GO; GO:0006548; P:viral-induced cell-cell fusion; IEA.
 DR InterPro; IPR000776; Fusion_gly.
 DR Pfam; PF00523; fusion_gly; MW; 609EE0247E59C54 CRC64;
 SQ SEQUENCE 550 AA; 59559 MW;

Query Match 45.0%; Score 72; DB 12; Length 550;
 Best Local Similarity 100.0%; Pred. No. 0.075; Indels 0; Gaps 0;

Qy 3 LSEIKGVIVHRLLEGV 17
 Db 288 LSEIKGVIVHRLLEGV 302

RESULT 15
 Q9QEW8 PRELIMINARY; PRT; 550 AA.
 ID Q9QEW8;
 AC Q9QEW8;
 DT 01-MAY-2000 (TREMBLrel. 13, Created)
 DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
 DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
 DE Fusion protein.
 OS Measles virus.
 OC ssRNA negative-strand viruses; Mononegavirales;
 OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.
 OC NCBI_TaxID=11234;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=OSA-2;
 RA Ning X., Ayata M., Morimoto K., Ito N., Shingai M., Kimura M., Ogura H.;
 RT "Nucleotide sequences of the fusion protein gene of subacute sclerosing panencephalitis viruses: deduced amino acid sequences showed the cytoplasmic domain highly mutated --truncated, elongated or predicted secondary structure changed.";
 RT Submitted (AUG-1999) to the EMBL/GenBank/DDBJ databases.
 RL EMBL; AF179437; AAF027021; -.
 DR PIR; PQ0376; PQ0376.
 DR HSSP; P04849; 1SVP.
 DR GO; GO:0019039; P:viral-cell fusion molecule activity; IEA.
 DR GO; GO:0006548; P:viral-induced cell-cell fusion; IEA.
 DR InterPro; IPR000776; Fusion_gly.
 DR Pfam; PF00523; fusion_gly; MW; 086EE51FED235BBB CRC64;
 SQ SEQUENCE 550 AA; 59315 MW;

Query Match 45.0%; Score 72; DB 12; Length 550;
 Best Local Similarity 100.0%; Pred. No. 0.075; Indels 0; Gaps 0;

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Qy 3 LSEIKGYIVHRLDEGV 17
Db 288 LSEIKGYIVHRLDEGV 302

Search completed: March 10, 2004, 09:25:31
Job time : 32.631 sees

PT or its analog.
 XX Claim 11; Page 7; 43pp; English.
 XX The invention relates to a synthetic immunogen for inducing specific antibodies against gonadotropin releasing hormone (GnRH) also known as luteinising hormone releasing hormone, (LHRH) comprising a fusion peptide which comprises a promiscuous helper T-cell peptide epitope and immunomimetic peptide epitope or its analogue. The synthetic immunogen is useful inducing an immune response against GnRH in an animal subject, and as such is useful as a contraceptive and in the treatment of diseases such as cancer (of the breast, uterus and other gynaecological cancer), endometriosis, uterine fibroids, benign prostatic hypertrophy and prostate cancer. The immunogen is effective in eliciting high and specific anti-GnRH antibody titres. The present sequence is a synthetic immunogen of the invention
 XX Sequence 31 AA;
 Query Match Score 159; DB 5; Length 31;
 Best Local Similarity 100.0%; Pred. No. 9.3e-17;
 Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 KLLSIKGVIVHLLEGVGPSPSLHMSYGLRP 30
 Db 1 KLLSIKGVIVHLLEGVGPSPSLHMSYGLRP 30

RESULT 2
 AAU11428 standard; peptide; 47 AA.
 ID AAU11428
 XN AAU11428;
 XX DT 12-MAR-2002 (first entry)
 XX DE Synthetic immunogen peptide 9.
 XX Gonadotrophin releasing hormone; GnRH; synthetic immunogen;
 KW Luteinising hormone releasing hormone; LHRH; contraceptive;
 KW promiscuous helper T-cell peptide epitope; immunomimetic Peptide epitope;
 KW breast cancer; uterine cancer; gynaecological cancer; endometriosis;
 KW uterine fibroid; benign prostatic hypertrophy; prostate cancer.
 OS Plasmodium falciparum.
 OS Mammalia.
 OS Synthetic.
 OS Chimeric.
 XX Location/Qualifiers
 PT Peptide 1..10 /note= "Gonadotrophin releasing hormone epitope (1..10 aa)"
 PT Misc-difference 1 /label= OTHER /note= "Other= Pyro-glutamic acid or 5-oxo proline"
 PT Peptide 11..16 /note= "Spacer peptide"
 PT Peptide 17..34 /note= "Malaria CSP protein (288-302 aa)"
 PT Peptide 35..38 /note= "Spacer peptide"
 PT Peptide 39..47 /note= "Gonadotrophin releasing hormone epitope (2-10 aa)"
 PT Modified-site 47 /note= "Amidated glycine or glycaminide"
 XX WO200185763-A2.
 PN PR 15-NOV-2001.
 PD XX PR 04-MAY-2001; 2001WO-US014363.

XX PR 05-MAY-2000; 2000US-020228P.
 XX PA (APHT-) APHTON CORP.
 XX PI Grimes S, Michaeli D, Stevens VC;
 XX DR WPI; 2002-049440/06.
 XX Novel synthetic immunogen for inducing immune response against gonadotropin releasing hormone, comprises fusion peptide having promiscuous helper T-cell peptide epitope and immunomimetic peptide epitope or its analog.
 XX Claim 11; Page 11; 43pp; English.
 XX The invention relates to a synthetic immunogen for inducing specific antibodies against gonadotropin releasing hormone (GnRH) also known as luteinising hormone releasing hormone, (LHRH) comprising a fusion peptide which comprises a promiscuous helper T-cell peptide epitope and immunomimetic peptide epitope or its analogue. The synthetic immunogen is useful inducing an immune response against GnRH in an animal subject, and as such is useful as a contraceptive and in the treatment of diseases such as cancer (of the breast, uterus and other gynaecological cancer), endometriosis, uterine fibroids, benign prostatic hypertrophy and prostate cancer. The immunogen is effective in eliciting high and specific anti-GnRH antibody titres. The present sequence is a synthetic immunogen of the invention
 XX Sequence 47 AA;
 Query Match Score 159; DB 5; Length 47;
 Best Local Similarity 100.0%; Pred. No. 1.5e-16;
 Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 KLLSIKGVIVHLLEGVGPSPSLHMSYGLRP 30
 Db 17 KLLSIKGVIVHLLEGVGPSPSLHMSYGLRP 46

RESULT 3
 AAR62705 standard; peptide; 25 AA.
 ID AAR62705
 XX AC AAR62705;
 XX DT 25-MAR-2003 (revised)
 DT 10-SEP-1995 (first entry)
 XX DE LHRH-containing immunogenic peptide.
 XX PH Helper T cell epitope; universal immune stimulator; invasin; hapten;
 KW vaccine; LHRH; luteinising hormone releasing hormone; prostate;
 KW androgen-dependent carcinoma; antitumour; infertility;
 KW measles virus F Protein.
 XX OS Synthetic.
 XX PH Key
 PT Domain 1..15 /note= "measles virus F protein helper T cell epitope"
 PT Domain 16..25 /note= "LHRH hapten"
 XX PN WO94250560-A1.
 XX PD 10-NOV-1994.
 XX PF 28-APR-1994;
 XX PR 27-APR-1993;
 PR 14-APR-1994;
 XX PP 94WO-US004832.
 XX PR 93US-00057166.
 PR 94US-0022975.

PA (LADD/) LADD A. E.
 PA (WANG/) WANG C. Y.
 PA (ZAMB/) ZAMB T.
 XX
 PI Ladd AE, Wang CY, Zamb T;
 XX
 DR WPI: 1994-357910/44.
 XX
 PT Immunogenic luteinising hormone releasing hormone peptide(s) - that
 suppress LHRH activity in males and females.
 XX
 P1 Ladd AE, Wang CY, Zamb T;
 P1
 P8 Claim 8; Page 84; 213pp; English.
 XX
 Synthetic immunogenic Peptides are provided in which a universal immune
 stimulator is linked to a peptide or protein hapten containing B cell
 and/or cytotoxic T lymphocyte epitopes, giving a product which causes
 potent immune responses to the coupled peptide or protein. The stimulator
 consists of (A) a promiscuous helper T cell epitope (Th) which elicits an
 immune response in members of a heterogeneous
 population expressing diverse HLA phenotypes, and (B) an adjuvant peptide
 sequence from the invasin protein of Yersinia. Spacer amino acid
 sequences (e.g. Gly-Gly) can be provided between the invasin and Th
 domains and between the immune stimulator and hapten components. When the
 hapten is LHRH, then optionally the invasin domain can be omitted from
 the immune stimulator component. The present sequence represents an LHRH-
 containing, invasin-free immunogenic peptide as above which can be used
 as a potent vaccine for treating e.g. prostatic hyperplasia, androgen-
 dependent carcinoma, prostatic carcinoma, testicular carcinoma,
 endometriosis, benign uterine tumours, recurrent functional ovarian
 cysts, (severe) premenstrual syndrome or oestrogen-dependent breast
 cancer, or for induction of infertility. (Updated on 25-MAR-2003 to
 correct PN field.)
 XX
 Sequence 25 AA;

Query Match 3 LSEIKGVIVHRLLEGVBPSLHNSYGLRP 30
 Best Local Similarity 73.1%; Score 117; DB 2; Length 25;
 Matches 24; Conservative 0; Mismatches 0; Indels 4; Gaps 1;

DB 1 LSEIKGVIVHRLLEGVBPSLHNSYGLRP 24

Qy 3 LSEIKGVIVHRLLEGVBPSLHNSYGLRP 30
 DB 1 LSEIKGVIVHRLLEGVBPSLHNSYGLRP 24

RESULT 4
 AAR62708
 ID AAR62708 standard; peptide; 27 AA.
 XX
 AC AAR62708;
 AC
 DT 25-MAR-2003 (revised)
 DT 10-SEP-1995 (first entry)
 DE LHRH-containing immunogenic peptide.
 XX
 Helper T cell epitope; universal immune stimulator; invasin; hapten;
 KW vaccine; LHRH; luteinising hormone releasing hormone; prostate;
 KW androgen-dependent carcinoma; antitumour; infertility;
 KW measles virus F protein.
 XX
 OS Synthetic.
 XX
 FH Location/Qualifiers
 FT 1..15
 /note= "measles virus F protein helper T cell epitope"
 FT 16..30
 /note= "measles virus F protein helper T cell epitope"
 FT 33..42
 /note= "LHRH hapten"
 FT Domain
 AX WO9425060-A1.
 XX 10-NOV-1994.

XX
 PA 28-APR-1994; 94WO-US004832.
 XX
 PR 27-APR-1993; 93US-0057166.
 PR 14-APR-1994; 94US-0023275.
 XX
 PA (LADD/) LADD A. E.
 PA (WANG/) WANG C. Y.
 PA (ZAMB/) ZAMB T.
 XX
 PI Ladd AE, Wang CY, Zamb T;
 XX
 WPI; 1994-357910/44.

XX
 DR
 PT Immunogenic luteinising hormone releasing hormone peptide(s) - that
 suppress LHRH activity in males and females.
 XX
 PS Claim 8; Page 86; 213pp; English.
 XX
 Synthetic immunogenic peptides are provided in which a universal immune
 stimulator is linked to a peptide or protein hapten containing B cell
 and/or cytotoxic T lymphocyte epitopes, giving a product which causes
 potent immune responses to the coupled peptide or protein. The stimulator
 consists of (A) a promiscuous helper T cell epitope (Th) which elicits an
 immune response to the coupled peptide in members of a heterogeneous
 population expressing diverse HLA phenotypes, and (B) an adjuvant peptide
 sequence from the invasin protein of Yersinia. Spacer amino
 acid sequences (e.g. Gly-Gly) can be provided between the invasin and Th
 domains and between the immune stimulator and hapten components. When the
 hapten is LHRH, then optionally the invasin domain can be omitted from
 the immune stimulator component. The present sequence represents an LHRH-
 containing, invasin-free immunogenic peptide as above which can be used
 as a potent vaccine for treating e.g. prostatic hyperplasia, androgen-
 dependent carcinoma, prostatic carcinoma, testicular carcinoma, androgen-
 dependent carcinomas, benign uterine tumours, recurrent functional ovarian
 cysts, (severe) premenstrual syndrome or oestrogen-dependent breast
 cancer, or for induction of infertility. (Updated on 25-MAR-2003 to
 correct PN field.)
 XX
 Sequence 42 AA;

Query Match 3 LSEIKGVIVHRLLEGVBPSLHNSYGLRP 30
 Best Local Similarity 73.1%; Score 117; DB 2; Length 42;
 Matches 24; Conservative 1; Mismatches 2; Indels 2; Gaps 1;

DB 15 VLSEIKGVIVHRLLEGVBPSLHNSYGLRP 41

RESULT 5
 AAR62707
 ID AAR62707 standard; peptide; 27 AA.
 XX
 AC AAR62707;
 XX
 DT 25-MAR-2003 (revised)
 DT 10-SEP-1995 (first entry)
 XX
 DE LHRH-containing immunogenic peptide.
 XX
 Helper T cell epitope; universal immune stimulator; invasin; hapten;
 KW vaccine; LHRH; luteinising hormone releasing hormone; prostate;
 KW androgen-dependent carcinoma; antitumour; infertility;
 KW measles virus F protein.
 XX
 OS Synthetic.
 XX
 FH Location/Qualifiers
 FT 1..15
 /note= "measles virus F protein helper T cell epitope"
 FT 16..30
 /note= "measles virus F protein helper T cell epitope"
 FT 33..42
 /note= "LHRH hapten"
 FT Domain
 AX
 FN WO9425060-A1.
 XX
 PT 1..15
 /note= "measles virus F protein helper T cell epitope"
 PT 16..30
 /note= "LHRH hapten"
 PT
 PT

CC containing immunogenic peptide as above which can be used as a potent
 CC vaccine for treating e.g. prostatic hyperplasia, androgen-dependent
 CC carcinoma, prostatic carcinoma, testicular carcinoma, endometriosis;
 CC benign uterine tumours, recurrent functional ovarian cysts, (severe)
 CC premenstrual syndrome or oestrogen-dependent breast cancer, or for
 CC induction of infertility. (Updated on 25-MAR-2003 to correct PN field.)
 XX

SQ Sequence 45 AA;

Query Match 72.5%; Score 116; DB 2; Length 45;
 Best Local Similarity 85.7%; Pred. No. 5e-10; Indels 2; Gaps 1;
 Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;

Qy 3 LSEIKGVIVHLEGVGEGPSLHWSYGLRP 30
 Db 19 LSEIKGVIVHLEGVGEGE-HWSYGLRP 44

PS Sequence 45 AA;

Query Match 72.5%; Score 116; DB 2; Length 45;
 Best Local Similarity 85.7%; Pred. No. 5e-10; Indels 2; Gaps 1;
 Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;

Qy 3 LSEIKGVIVHLEGVGEGPSLHWSYGLRP 30
 Db 19 LSEIKGVIVHLEGVGEGE-HWSYGLRP 44

PS Sequence 45 AA;

Query Match 72.5%; Score 116; DB 2; Length 45;
 Best Local Similarity 85.7%; Pred. No. 5e-10; Indels 2; Gaps 1;

Qy 3 LSEIKGVIVHLEGVGEGPSLHWSYGLRP 30
 Db 19 LSEIKGVIVHLEGVGEGE-HWSYGLRP 44

PS Sequence 45 AA;

Qy 3 LSEIKGVIVHLEGVGEGPSLHWSYGLRP 30
 Db 19 LSEIKGVIVHLEGVGEGE-HWSYGLRP 44

RESULT 10
 ID AAY91163 Standard; peptide; 27 AA.
 XX
 AC AAY91163;
 XX
 DT 12-SEP-2003 (revised)
 DT 22-MAY-2000 (first entry)

XX
 DE Modified MVF Th epitope/LHRH antigenic peptide, SEQ ID NO:43.
 XX
 KW Promiscuous T-cell epitope; measles virus F Protein; MVF;
 KW hepatitis B virus surface antigen; HBV; immunogenic; B-cell epitope;
 KW luteinising hormone releasing hormone; LHRH; contraceptive; anticancer;
 KW bonatocatin; growth promotion; CD4 receptor; HIV-1; antiviral; FMDV;
 KW foot and mouth disease virus; immunoglobulin B; IgB; anti-allergic;
 KW Plasmocitum falciiparum; circumsporozoite; antimalarial; CETP;
 KW cholestryl ester transport protein; anti-arteriosclerotic.
 XX
 OS Measles virus.
 OS Rattus sp.
 OS Chimeric.
 XX
 PN WO966957-A2.
 XX
 PD 29-DEC-1999.
 XX
 PP 21-JUN-1999; 99WO-US013975.
 XX
 PR 20-JUN-1998; 98US-0010412.
 XX
 PA (UNBI-) UNITED BIOMEDICAL INC.
 XX
 PI Wang CY;
 DR 2000-16564/14.
 XX
 PT New artificial T helper cell epitope and derived immunogens with target
 PT antigenic site, for immunization against e.g. malaria, arteriosclerosis
 PT or human immune deficiency virus.
 XX
 PS Example 1; Page 80; 129pp; English.

XX
 CC The invention relates to novel promiscuous T helper cell epitopes (Th),
 CC and immunogenic peptides comprising the Th epitopes of the invention
 CC along with B cell epitopes. The Th epitopes and peptide immunogens
 CC containing them, are used to induce a T helper cell response,
 CC specifically against Plasmodium falciparum, cholesterol ester transport
 CC protein (CETP) or HIV epitopes, but more generally against any pathogen,
 CC immunoreactive self-antigen or tumour antigen. The Th epitopes and
 CC peptide immunogens may be used for prevention and/or treatment of
 CC infections (HIV, foot-and-mouth disease or malaria); for cancer
 CC immunotherapy, for inhibition of the action of luteinising hormone
 CC releasing hormone (LHRH) for contraception, treatment of boar taint, in meat, and immunocastration
 CC dependent cancer, prevention of the growth of animals; or for treating allergies or
 CC arteriosclerosis. Incorporation of a promiscuous Th (functional in
 CC genetically diverse subjects) into an immunogen improves capacity to
 CC induce a strong T helper cell-mediated immune response, resulting in
 CC production of antibodies against a target antigen. Th can replace carrier
 CC proteins and pathogen-derived T helper epitopes. Sequence AA91121
 CC represents a promiscuous T helper epitope from the measles virus F (MVF)
 CC protein and sequence AAY91122-Y91142, AA91126 and AAY91245-Y91246
 CC represent synthetic Th epitopes based on the MVF Th epitope. Sequence
 CC AAY91143 represents a promiscuous Th epitope from hepatitis B virus (HBV)
 CC surface antigen, and sequences AAY91144-Y91155 are synthetic epitopes
 CC derived from this HBV epitope. AAY91156-Y91196, AA91127 and AA91242-
 CC derived from this HBV epitope.

Query Match 72.5%; Score 116; DB 7; Length 45;
 Best Local Similarity 85.7%; Pred. No. 5e-10; Indels 2; Gaps 1;

SQ Sequence 45 AA;

Y91244 are antigenic peptides comprising an LHRH sequence joined to a promiscuous Th epitope. AAY9197 is the LHRH target antigenic peptide used in these LHRH antigenic peptides. AAY91200 is somatostatin, and AAY91201-Y91207 are antigenic peptides comprising somatostatin and a Th epitope. Somatostatin immunogens may be used to promote growth in livestock. AAY91208 is a human CD4 CDR2-like domain antigenic site, and AAY91209-Y90211 are MNH Th epitope/CD4 CDR2 antigenic peptides which may be used to prevent HIV infection of T cell lines. AAY90212 is a modified version of a human IgE (immunoglobulin E) CH3 domain, and AAY0213-Y90219 are Th epitope/IgE CH3 antigenic peptides which may be used in the treatment of allergies. AAY9120 is a peptide derived from foot and mouth disease virus (FMDV) VP1 capsid protein and AAY91221-Y91222 comprise this peptide and a Th epitope. AAY91223 is a Plasmodium falciparum circumsporozoite (CS) target antigen, and AAY9124-Y91225 comprise the CS antigen and an MVE Th epitope and may be used in a malaria vaccine. AAY91228-Y91231 represent CETP-derived Peptides and AAY91232-Y91241 are immunogens comprising a CETP peptide and a Th epitope which may be used to prevent or treat arteriosclerosis and cardiovascular disease. AAY91247 and AAY91252-Y91257 are HIV-1 neutralising B-cell epitopes, and AAY91248-Y91251 and AAY91258-Y91273 are antigenic peptides comprising MWH Th and HIV-1 B-cell epitope which may be used as a component in an anti-HIV-1 vaccine. AAY9118 and AAY91199 are respectively an immunostimulatory invasin protein epitope from Yersinia species, and hinge spacer Peptide, both of which may optionally be used in the antigenic peptides of the invention. (Updated on 12-SEP-2003 to standardise OS field)

Sequence 27. AA:

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Query Match          70.6%; Score 113; DB 3; length 27;
Best Local Similarity 82.1%; Pred. No. 7.e-10;
Matches      23; Conservative 1; Mismatches 2; Indels 0
3 LSETKGVIVHLRVEGEGPSLHWSKGIRP 30

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1 LSBTKGVTVHKGEGCGE--HWSYGLRP 26

AXY91175,
12-SEP-2003 (revised)
22-MAY-2000 (first entry)

Modified MVF Th epitope/LHRH antigenic peptide, SEQ ID NO:55.

Promiscuous T-cell epitope; measles virus F protein; MVF;
hepatitis B virus surface antigen; HBV; immunogenic; B-cell e
luteinising hormone releasing hormone; LHRH; contraceptive;
sonatastatin; growth promotion; CD4 receptor; HIV 1; antivira
foot and mouth disease virus; immunoglobulin B; IgE; anti-all
Plasmodium falciparum; circumsPorozoite; antimarial; CEP7;
cholasteryl ester transport protein; anti-arteriosclerotic.

Measles virus.
RATTUS SD.

Chimeric.

WO966957-A2.
29-DEC-1999.
21-JUN-1999; 99WO-US013975.
20-JUN-1998; 98US-00100412.
(UNIBI-) UNITED BIOMEDICAL INC.

Wang CY;
WPI: 2000-160554/14

XX New artificial T helper cell epitope and derived immunogens with target antigen site, for immunization against e.g. malaria, arteriosclerosis

Example 1; Page 84; 129pp; English.

The invention relates to novel promiscuous T helper cell epitopes (Th_1) and immunogenic peptides comprising the Th_1 epitopes of the invention along with B cell epitopes. The Th_1 epitopes and peptide immunogens containing them, are used to induce a T helper cell response, specifically against Plasmodium falciparum, cholesteryl ester transport protein (CEBP) or HIV epitopes, but more generally against any pathogen immunoreactive self-antigen or tumour antigen. The Th_1 epitopes and peptide immunogens may be used for prevention and/or treatment of infections (HIV, foot-and-mouth disease or malaria); for cancer immunotherapy; for inhibition of the action of luteinising hormone-releasing hormone (LHRH) for contraception; treatment of hormone-dependent cancers; prevention of boar taint in meat; and immunocastration; for promoting the growth of animals; or for treating allergies or arteriosclerosis. Incorporation of a promiscuous Th_1 functional in genetically diverse subjects into an immunogen improves capacity to induce a strong T helper cell-mediated immune response, resulting in production of antibodies against a target antigen. This can replace carrier proteins and pathogen-derived T helper epitopes. Sequence AAY91212 represents a promiscuous T helper epitope from the measles virus F (MV) genome.

protein and sequences AAY91122-Y91142, AAY91226 and AY91245-Y91246 represent synthetic Th epitopes based on the MVE Th epitope. Sequence AAY91143 represents a promiscuous Th epitope from hepatitis B virus (HBV) surface antigen, and sequences AAY91144-Y91155 are synthetic epitopes derived from this HBV epitope. AAY91156-Y91196, AAY911227 and AAY91244 are antigenic peptides comprising an LHRH sequence joined to a promiscuous Th epitope. AAY91197 is the LHRH target antigenic peptide used in these LHRH antigenic peptides. AAY91200 is somatostatin, and AAY91201-Y91207 are antigenic peptides comprising somatostatin and a Th epitope. Somatostatin immunogens may be used to promote growth in livestock. AAY91208 is a human CD4 CDR2-like domain antigenic site, and AAY91209-Y90111 are MVE Th epitope/CD4 CDR2 antigenic peptides which may be used to prevent MVE infection of T cells. AY90112 is a modified version of a human IgE (immunoglobulin E) CH3 domain, and AAY9023-Y9024 are Th epitope/IgE CH3 antigenic peptides which may be used in the treatment of allergies. AAY91220 is a peptide derived from foot and mouth disease virus (FMDV) VP1 capsid protein and AAY91221-Y91222 comprise the peptide and a Th epitope. AAY91223 is a Plasmodium falciparum circumsporozoite (CS) target antigen, and AAY91224-Y91225 comprise the antigen and an MVE Th epitope and may be used in a malaria vaccine. AAY91228-Y91231 represent CEBP-derived peptides and AAY91232-Y91241 are immunogens comprising a CEBP peptide and a Th epitope which may be used to prevent or treat arteriosclerosis and cardiovascular disease. AAY91229 and AAY91252-Y91257 are HIV-1 neutralising B-cell epitopes, and AAY91244-Y91251 and AAY91258-Y91273 are antigenic peptides comprising MVH TH and HIV-1 B-cell epitope which may be used as a component in an anti-HIV-1 vaccine. AAY91198 and AAY91199 are respectively an immunomimulatory invasin protein epitope from Yersinia species, and hinge spacer peptides which may optionally be used in the antigenic peptides of the

CC invention. (Updated on 12-SEP-2003 to standardise OS field)
 XX
 SQ Sequence 31 AA:
 Query Match 68.8%; Score 110; DB 3; Length 31;
 Best Local Similarity 78.6%; Pred. No. 2.6e-09;
 Matches 22; Conservative 1; Mismatches 5; Indels 0; Gaps 0
 Qy 3 LSEIKGVIVKHLLEGVYEGPSLHWSYGLRP 30
 | : ||| : ||| : ||| : ||| : ||| : ||| : ||| : |||
 Db 3 LSEIKGVIVKHLLEGVYEGPSLHWSYGLRP 30

RESULT 12
AY91161
IDD AY91161 standard; peptide; 27 AA.
XXX AC AY91161;

XX 12-SEP-2003 (revised)
 DT 22 MAY 2000 (first entry)

XX Modified MVF Th epitope/LHRH antigenic peptide, SEQ ID NO:41.

XX Promiscuous T-cell epitope; measles virus F protein; MVE;
 hepatitis B virus surface antigen; HBV; immunogenic; B-cell epitope;
 luteinising hormone releasing hormone; LHRH; contraceptive; anticancer;
 somatoatin; growth promotion; CD4 receptor; HIV-1; antiviral; FMDV;
 foot and mouth disease virus; immunoglobulin E; IgE; anti-allergic;
 Plasmodium falciparum; circumsporozoite; antimalarial; CEPF;
 cholesterol ester transport protein; anti-arteriosclerotic.

XX Measles virus.
 OS Rattus sp.
 OS Chimeric.

XX WO9966957-A2.
 XX PD 29-DEC-1999.
 XX PF 21-JUN-1999; 99WO-US013975.
 XX PR 20-JUN-1998; 98US-00100412.
 XX PA (UNBI-) UNITED BIOMEDICAL INC.

XX PI Wang CY;
 XX DR 2000-160564/14.
 XX PT New artificial T helper cell epitope and derived immunogens with target
 PT antigenic site, for immunization against e.g. malaria, arteriosclerosis
 PT or human immune deficiency virus.

PS Example 1; Page 79; 129PP; English.

XX The invention relates to novel promiscuous T helper cell epitopes (Th),
 CC and immunogenic peptides comprising the Th epitopes of the invention
 along with B cell epitopes. The Th epitopes and peptide immunogens
 containing them, are used to induce a T helper cell response,
 specifically against Plasmodium falciparum, cholestrylo ester transport
 protein (CEP) or HIV epitopes, but more generally against any pathogen,
 CC immunoreactive self-antigen or tumour antigen. The Th epitopes and
 peptide immunogens may be used for prevention and/or treatment of
 CC infections (HTV, foot-and-mouth disease or malaria); for cancer
 immunotherapy; for inhibition of the action of luteinising hormone
 CC releasing hormone (LHRH) for contraception, treatment of hormone-
 dependent cancer, prevention of boar taint in meat, and immunocastration
 CC ; for promoting the growth of animals; or for treating allergies or
 arteriosclerosis. Incorporation of a promiscuous Th (functional in
 CC genetically diverse subjects) into an immunogen improves capacity to
 induce a strong T helper cell-mediated immune response, resulting in
 CC production of antibodies against a target antigen. Th can replace carrier
 CC proteins and pathogen-derived T helper epitopes. Sequence AAY91121
 CC represents a promiscuous T helper epitope from the measles virus F (MVF)
 CC protein and sequences AAY91122-Y91142, AAY91226 and AAY91245-Y91246
 CC represent synthetic Th epitopes based on the MVF Th epitope. Sequence
 CC AAY91143 represents a promiscuous Th epitope from hepatitis B virus (HBV)
 CC surface antigen, and sequences AAY91144-Y91155 are synthetic epitopes
 CC derived from this HBV epitope. AAY91156-Y91196, AAY91227 and AAY91142-
 Y91244 are antigenic peptides comprising an LHRH sequence joined to a
 CC promiscuous Th epitope. AAY91197 is the LHRH target antigenic peptide
 CC used in these LHRH antigenic peptides. AAY91200 is somatosatin, and
 CC AAY91201-Y91207 are antigenic peptides comprising somatosatin and a Th
 CC epitope. Somatotropin immunogens may be used to promote growth in
 CC livestock. AAY91208 is a human CD4 CDR2-like domain antigenic site, and
 CC AAY9109-Y90211 are MVH Th epitope/CD4 CDR2 antigenic peptides which may
 CC be used to prevent HIV infection of T cells. AAY90212 is a modified
 CC version of a human IgE (immunoglobulin E) CH3 domain, and AAY90213-Y90219
 CC are Th epitope/IGE CH3 antigenic peptides which may be used in the
 CC treatment of allergies. AAY91220 is a peptide derived from foot and mouth

CC disease virus (FMDV) VP1 capsid protein and AAY91221-Y91222 comprise this
 CC peptide and a Th epitope. AAY91223 is a Plasmodium falciparum
 CC circumsporozoite (CS) target antigen, and AAY91224 Y91225 comprise the CS
 CC antigen and an MVH Th epitope and may be used in a malaria vaccine.
 CC AAY91228-Y91231 represent CEPF-derived peptides and AAY91232-Y91241 are
 CC immunogens comprising a CEPF peptide and a Th epitope which may be used
 CC to prevent or treat arteriosclerotic and cardiovascular disease. AAY91247
 CC and AAY91252-Y91257 are HIV-1 neutralising B-cell epitopes, and AAY91248-
 CC Y91251 and AAY91258-Y91273 are antigenic peptides comprising MVH Th and
 CC HIV-1 B-cell epitope which may be used as a component in an anti-HIV-1
 CC vaccine. AAY9198 and AAY9199 are respectively an immunostimulatory
 CC invasin protein epitope from Yersinia species, and hinge spacer peptides
 CC both of which may optionally be used in the antigenic peptides of the
 CC invention. (updated on 12-SEP-2003 to standardise OS field)

XX OS Sequence 27 AA:
 XX SQ Rattus sp.
 XX Query Match 67.5%; Score 108; DB 3; Length 27;
 XX Best Local Similarity 71.4%; Prod. No. 4.4e-09;
 XX Matches 20; Conservative 4; Mismatches 2; Indels 2; Gaps 1;

XX CY 3 LSEIKGVIVHLRLEGEGPSLHWYSGLRP 30
 XX Db 1 ISEIKGVIVHLRLEGEGPSLHWYSGLRP 26

RESULT 13
 AAY91167
 ID AAY91167 standard; peptide; 27 AA.
 XX AC AAY91167;
 XX DE Modified MVF Th epitope/LHRH antigenic peptide, SEQ ID NO:47.
 XX DT 12-SEP-2003 (revised)
 XX DT 22-MAY-2000 (first entry)

XX DE Modified MVF Th epitope/LHRH antigenic peptide, SEQ ID NO:47.
 XX AC AAY91167;
 XX DE Promiscuous T-cell epitope; measles virus F protein; MVE;
 XX KW hepatitis B virus surface antigen; HBV; immunogenic; B-cell epitope;
 XX KW luteinising hormone releasing hormone; LHRH; contraceptive; anticancer;
 XX KW somatostatin; growth promotion; CD4 receptor; HIV-1; antiviral;
 XX KW foot and mouth disease virus; immunoglobulin E; IgE; anti-allergic;
 XX KW Plasmodium falciparum; circumsporozoite; antimalarial; CEPF;
 XX KW cholestrylo ester transport protein; anti-arteriosclerotic.
 XX Measles virus.
 XX OS Rattus sp.
 XX OS Chimeric.

XX PN WO966957-A2.
 XX PR 29-DEC-1999.
 XX PD 29-DEC-1999.
 XX PP 21-JUN-1999; 99WO-US013975.
 XX PR 20-JUN-1998; 98US-00100412.
 XX PR (UNBI-) UNITED BIOMEDICAL INC.
 XX PA Wang CY;
 XX DR 2000-160564/14.
 XX PT New artificial T helper cell epitope and derived immunogens with target
 PT antigenic site, for immunization against e.g. malaria, arteriosclerosis
 PT or human immune deficiency virus.

PS Example 1; Page 81; 129PP; English.

XX The invention relates to novel promiscuous T helper cell epitopes (Th),
 CC and immunogenic peptides comprising the Th epitopes of the invention
 along with B cell epitopes. The Th epitopes and peptide immunogens
 containing them, are used to induce a T helper cell response,
 specifically against Plasmodium falciparum, cholestrylo ester transport
 protein (CEP) or HIV epitopes, but more generally against any pathogen,
 CC immunoreactive self-antigen or tumour antigen. The Th epitopes and
 peptide immunogens may be used for prevention and/or treatment of
 CC infections (HTV, foot-and-mouth disease or malaria); for cancer
 immunotherapy; for inhibition of the action of luteinising hormone
 CC releasing hormone (LHRH) for contraception, treatment of hormone-
 dependent cancer, prevention of boar taint in meat, and immunocastration
 CC ; for promoting the growth of animals; or for treating allergies or
 arteriosclerosis. Incorporation of a promiscuous Th (functional in
 CC genetically diverse subjects) into an immunogen improves capacity to
 induce a strong T helper cell-mediated immune response, resulting in
 CC production of antibodies against a target antigen. Th can replace carrier
 CC proteins and pathogen-derived T helper epitopes. Sequence AAY91121
 CC represents a promiscuous T helper epitope from the measles virus F (MVF)
 CC protein and sequences AAY91122-Y91142, AAY91226 and AAY91245-Y91246
 CC represent synthetic Th epitopes based on the MVF Th epitope. Sequence
 CC AAY91143 represents a promiscuous Th epitope from hepatitis B virus (HBV)
 CC surface antigen, and sequences AAY91144-Y91155 are synthetic epitopes
 CC derived from this HBV epitope. AAY91156-Y91196, AAY91227 and AAY91142-
 Y91244 are antigenic peptides comprising an LHRH sequence joined to a
 CC promiscuous Th epitope. AAY91197 is the LHRH target antigenic peptide
 CC used in these LHRH antigenic peptides. AAY91200 is somatosatin, and
 CC AAY91201-Y91207 are antigenic peptides comprising somatosatin and a Th
 CC epitope. Somatotropin immunogens may be used to promote growth in
 CC livestock. AAY91208 is a human CD4 CDR2-like domain antigenic site, and
 CC AAY9109-Y90211 are MVH Th epitope/CD4 CDR2 antigenic peptides which may
 CC be used to prevent HIV infection of T cells. AAY90212 is a modified
 CC version of a human IgE (immunoglobulin E) CH3 domain, and AAY90213-Y90219
 CC are Th epitope/IGE CH3 antigenic peptides which may be used in the
 CC treatment of allergies. AAY91220 is a peptide derived from foot and mouth

specifically against Plasmodium falciparum, cholesteryl ester transport protein (CETP) or HIV epitopes, but more generally against any pathogen, immunoreactive self-antigen or tumour antigen. The Th epitopes and peptide immunogens may be used for prevention and/or treatment of infections (HIV, foot-and-mouth disease or malaria); for cancer immunotherapy; for inhibition of the action of luteinising hormone-releasing hormone (LHRH) for contraception, treatment of hormone-dependent cancer, prevention of boar taint in meat, and immunocastration for promoting the growth of animals; or for treating allergies or arterioclerosis. Incorporation of a promiscuous Th (functional in genetically diverse subjects) into an immunogen improves capacity to induce a strong T helper cell-mediated immune response, resulting in production of antibodies against a target antigen. Th can replace carrier proteins and pathogen-derived T helper epitopes. Sequence AAY9112-1 represents a promiscuous T helper epitope from the measles virus P (MV) protein and sequences AAY9112-2-Y91142, AAY91226 and AAY91245-Y91246 represent synthetic Th epitopes based on the MV Th epitope. Sequence AAY91143 represents a promiscuous Th epitope from hepatitis B virus (HBV surface antigen, and sequences AAY91144-Y91155 are synthetic epitopes derived from this HBV epitope. Sequence AAY9116-Y91196, AAY91227 and AAY91244 are antigenic peptides comprising an LRRH sequence joined to a promiscuous Th epitope. AAY91197 is the LRRH target antigenic peptide used in these LRRH antigenic peptides. AAY91200 is somatostatin, and AAY91201-Y91207 are antigenic peptides comprising somatostatin and a Th epitope. Somatostatin immunogens may be used to promote growth in livestock. AAY91208 is a human CD4 CDR2-like domain antigenic site, and AAY91209-Y90211 are MVA Th epitope/CD4 CDR2 antigenic Peptides which may be used to prevent HIV infection of T cells. AAY90212 is a modified version of a human IgE (immunoglobulin E) CH3 domain, and AAY90213-Y90214 are Th epitope/IgE CH3 antigenic Peptides which may be used in the treatment of allergies. AAY91220 is a peptide derived from foot and mouth disease virus (FMDV) VP1 capsid protein and AAY91221-Y91222 comprise the C peptide and a Th epitope. AAY91223 is a Plasmid falciparum circumsporozoite (CS) target antigen, and AAY91224-Y91225 comprise the CS antigen and an MVA Th epitope and may be used in a malaria vaccine. AAY91228-Y91231 represent CETP-derived peptides and AAY91232-Y91241 are immunogens comprising a CETP peptide and a Th epitope which may be used to prevent or treat arteriosclerosis and cardiovascular disease. AAY91224 and AAY91225-Y91257 are HIV-1 neutralising B-cell epitopes, and AAY91248-Y91251 and AAY91258-Y91273 are antigenic peptides comprising MVA Th and HIV-1 B-cell epitope which may be used as a component in an anti-HIV-1 vaccine. AAY91198 and AAY91199 are respectively an immunostimulatory invasin protein epitope from Yersinia species, and hinge spacer peptide, both of which may optionally be used in the antigenic peptides of the invention. (Updated on 12-SEP-2003 to standardise OS field)

X	Q	Sequence 27 AA:	Score 108;	DB 3;	Length 27;	
		Query Match	67.5%	Pred. No. 4.4e-09;		
		Best Local Similarity	75.0%			
		Matchers 21; Conservative Matchers 21;		3; Mismatches 3;	Indels 2;	Gaps 2;
Y	:	LSEIKGIVHRLLEGVGPSTLHWSYGLRP	3.0			

XX	Synthetic.
OS	Yersinia sp.
OS	Measles virus.
OS	Unidentified.
XX	
PH	Key Peptide
PT	Peptide
PT	Peptide
PT	Peptide
PT	Peptide
PT	Peptide
PT	Peptide
PT	Peptide
PT	Peptide
PN	WO9966952-A1.
XX	
PD	29-DEC-1999.
XX	
PF	21-JUN-1999;
XX	
PR	20-JUN-1998;
XX	
PA	(UNB1-) UNITED
XX	
P1	Wang CY;
XX	
DR	WPI: 2000-16056
XX	
PT	New Peptide immuno-
PR	antigen site an-
PR	treatment of ca-
XX	
PS	Claim 9: Page 7
XX	The present seq-
CC	domain immunot-
CC	cell (Th) epitope
CC	hormone (LHRH).
CC	synthetic anti-
CC	modeled after
CC	Measles virus.

Journal of Health Politics, Policy and Law, Vol. 29, No. 4, December 2004
DOI 10.1215/03616878-29-4 © 2004 by The University of Chicago

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RP

odium falcipes, but tumour or smooth disease. The all-mediators against a target peptide. AAY91197 is a helper epitope. Y91197 is a specific peptide antigen. Peptides bases amino acids AA9119223 is a set antigen. It is a C-terminal peptide derivative of immunoglobulinic pentapeptide. Y91195 are from Vertebrates. It can be used from SEP-2003 to

— 2 —

11

specifically CEBTP protein (CEBTP) can induce immunoreactive substances (HABA) in the tumor tissue. The substances have cytotoxicity and can induce apoptosis in tumor cells. The mechanism of action of HABA is not clear. It may be due to the inhibition of the synthesis of DNA and RNA, or it may be due to the inhibition of the synthesis of proteins. The mechanism of action of HABA is not clear. It may be due to the inhibition of the synthesis of DNA and RNA, or it may be due to the inhibition of the synthesis of proteins.

KAY

RESULT	14
D	RAY68573
XX	RAY68573 standard; peptide; 45 AA.
AC	RAY68573;
CX	05-MAY-2000 (first entry)
DE	Peptide immunogen comprising a Th epitope and LHRH target antigen.
DX	Helper T cell epitope; F protein; Measles virus; peptide immunogen; LHRH luteinising hormone-releasing hormone; spermatogenesis; ovulation; oestrus; sexual development; sex hormone; promiscuous T helper epitope; vaccine; contraceptive; hormone-dependent tumour; Prostate Cancer; breast Cancer; endometriosis; boar taint; meat quality; invasin domain; immunocastration.

AA	Sequence 45 AA;	Score	DB 3;	Length	45;
SQ					
Query Match	67.5%;	Score	108;	DB 3;	Length
Best Local Similarity	71.4%;	Pred.	No.	8.2e-09;	45;
Matches 20;	Conservative 4;	Mismatches	2;	Indels	2;
Gaps					
Qy	3 LSEIKGVTVHRLLEGPGPSLHNSYGLRP	30			
Db	19 ISEIKGVTVHKGEGIGSE-HNSYGLRP	44			
RESULT 15					
AY91165					
ID AY91165	standard; peptide	45 AA.			
XX					
AC					
XX					

DT 12-SEP-2003 (revised)
DT 22-MAY-2000 (first entry)

Modified MVF Th epitope/LHRH antigenic peptide, SEQ ID NO:45.

DE Promiscuous T-cell epitope; measles virus F protein; MVF;
KW hepatitis B virus surface antigen; HBV; immunogenic; B-cell epitope;
KW luteinising hormone-releasing hormone; LHRH; contraceptive; anticancer;
KW somatotatin; growth promotion; CD4 receptor; HIV-1; antiviral; FMDV;
KW foot and mouth disease virus; immunoglobulin E; IgB; anti-allergic;
KW Plasmodium falciparum; circumsporozoite; anti-malarial; CETP;
KW cholesteroyl ester transport protein; anti-arteriosclerotic.

XX Measles virus.
OS Rattus sp.
OS Chimeric.

XX PN WO9966957-A2.

XX PD 29-DEC-1999.

XX PF 21-JUN-1999; 99WO-US013975.

XX PR 20-JUN-1998; 98US5-00100412.

XX PA (UNIBI-) UNITED BIOMEDICAL INC.

XX PI Wang CY;

XX DR 2000-160564/14.

XX PT New artificial T helper cell epitope and derived immunogens with target antigenic site, for immunization against e.g. malaria, arteriosclerosis or human immune deficiency virus.

XX PS Example 1: Page 80; 129PP; English.

XX The invention relates to novel promiscuous T helper cell epitopes (Th), and immunogenic peptides comprising the Th epitopes of the invention, along with B cell epitopes. The Th epitopes and peptide immunogens containing them, are used to induce a T helper cell response, specifically against Plasmodium falciparum, cholesteroyl ester transport protein (CETP) or HIV epitopes, but more generally against any pathogen, immunoreactive self-antigen or tumour antigen. The Th epitopes and peptide immunogens may be used for prevention and/or treatment of infections (HIV, foot-and-mouth disease or malaria); for cancer immunotherapy; for inhibition of the action of luteinising hormone-releasing hormone (LHRH) for contraception, treatment of hormone-dependent cancer, prevention of boar taint in meat, and immunocastration; for promoting the growth of animals; or for treating allergies or arteriosclerosis. Incorporation of a promiscuous Th (functional in genetically diverse subjects) into an immunogen improves capacity to induce a strong T helper cell-mediated immune response, resulting in production of antibodies against a target antigen. It can replace carrier proteins and pathogen-derived T helper epitopes. Sequence AAY91211 represents a promiscuous T helper epitope from the measles virus F (MVF) protein and sequences AAY91222-Y91142, AAY91226 and AAY91245-Y91246.

XX represent synthetic Th epitopes based on the MVF Th epitope. Sequence AAY91243 represents a promiscuous Th epitope from hepatitis B virus (HBV) surface antigen, and sequences AAY91144-Y91155 are synthetic epitopes derived from this HBV epitope. AAY91156-Y91196, AAY91227 and AAY91244 are antigenic peptides comprising an LHRH sequence joined to a promiscuous Th epitope. AAY91197 is the LHRH target antigenic peptide used in these LHRH antigenic peptides. AAY91200 is somatostatin, and AAY91201-Y91207 are antigenic peptides comprising somatostatin and a Th epitope. Somatostatin immunogens may be used to promote growth in livestock. AAY91208 is a human CD4 CDR2-like domain antigenic site, and AAY91109-Y90211 are MVH Th epitope/CD4 CDR2-like domain antigenic site, and CC both of which may optionally be used in the infection of T cells. AAY90212 is a modified version of a human IgE (immunoglobulin E) CH3 domain, and AAY0213-Y90219 are Th epitope/IgE CH3 antigenic peptides which may be used in the treatment of allergies. AAY91220 is a peptide derived from foot and mouth disease virus (FMDV) VP1 capsid protein and AAY91221-Y91222 comprise this invention. (Updated on 12-SEP-2003 to standardise OS field)

XX Sequence 45 AA;

SQ Query Match 67.5%; Score 108; DB 3; Length 45;
Best Local Similarity 71.4%; Prd. No. 8.2e-09;
Matches 20; Conservative 4; Mismatches 2; Indels 2; Caps 1;

Qy 3 LSEIKGVIVHLRLEGPGPSLAVSYGLRP 30
:|||||:|||:||:||:|||:|||:|||:
Db 19 ISEIKGVIVHLKIGGEE-HNSYGLRP 44

Search completed: March 10, 2004, 09:12:09
Job time : 47.6809 secs

GanCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using SW model

Run on: March 10, 2004, 09:16:59 ; Search time 4.71595 Seconds
(without alignments)
268.645 Million cell updates/sec

Title: US-09-848-834A-6

Perfect Score: 29

Sequence: 1 SSGSGL 6

Scoring table: BLOSSUM62 Gapext 10.0 , Gapext 0.5

Searched: 809742 seqs., 21153259 residues

Total number of hits satisfying chosen parameters: 809742

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 100%
Maximum Match 100%
Listing first 45 summariesDatabase : Published Applications AA:
1: /cgns_6/ptodata/2/pbpaas/US07_PUBCOMB.pep:
2: /cgns_6/ptodata/2/pbpaas/US07_PUBCOMB.pep:
3: /cgns_6/ptodata/2/pbpaas/US06_NEWPUB.pep:
4: /cgns_6/ptodata/2/pbpaas/US06_PUBCOMB.pep:
5: /cgns_6/ptodata/2/pbpaas/US07_NEWPUB.pep:
6: /cgns_6/ptodata/2/pbpaas/US08_PUBCOMB.pep:
7: /cgns_6/ptodata/2/pbpaas/US08_NEWPUB.pep:
8: /cgns_6/ptodata/2/pbpaas/US08_PUBCOMB.pep:
9: /cgns_6/ptodata/2/pbpaas/US09A_PUBCOMB.pep:
10: /cgns_6/ptodata/2/pbpaas/US09_PUBCOMB.pep:
11: /cgns_6/ptodata/2/pbpaas/US09C_PUBCOMB.pep:
12: /cgns_6/ptodata/2/pbpaas/US07_NEWPUB.pep:
13: /cgns_6/ptodata/2/pbpaas/US08_PUBCOMB.pep:
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15: /cgns_6/ptodata/2/pbpaas/US10_PUBCOMB.pep:
16: /cgns_6/ptodata/2/pbpaas/US10_NEWPUB.pep:
17: /cgns_6/ptodata/2/pbpaas/US60_NEWPUB.pep:
18: /cgns_6/ptodata/2/pbpaas/US60_PUBCOMB.pep:
;

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match	Length	DB ID	Description
1	29	100.0	6	9	US-09-848-834A-6	Sequence 6, Appli
2	29	100.0	8	9	US-09-848-834A-7	Sequence 7, Appli
3	29	100.0	31	9	US-09-848-834A-15	Sequence 15, Appli
4	29	100.0	34	9	US-09-848-834A-13	Sequence 13, Appli
5	29	100.0	36	9	US-09-848-834A-16	Sequence 16, Appli
6	29	100.0	37	9	US-09-848-834A-14	Sequence 14, Appli
7	29	100.0	46	9	US-09-848-834A-19	Sequence 19, Appli
8	29	100.0	47	9	US-09-848-834A-17	Sequence 17, Appli
9	29	100.0	50	9	US-09-848-834A-18	Sequence 18, Appli
10	29	100.0	51	9	US-09-848-834A-20	Sequence 20, Appli
11	29	100.0	53	9	US-09-848-761-47649	Sequence 47649, Appli
12	29	100.0	75	9	US-09-751-872-425	Sequence 425, Appli
13	29	100.0	75	10	US-09-776-937-425	Sequence 425, Appli
14	29	100.0	200	9	US-09-810-756-38	Sequence 38, Appli
15	29	100.0	200	9	US-09-874-585B-38	Sequence 38, Appli

Result No.	Score	Query	Match	Length	DB ID	Description	
16	29	100.0	29	100.0	US-09-731-626-6493	Sequence 6493, Appli	
17	29	100.0	29	100.0	US-10-210-110-34	Sequence 34, Appli	
18	29	100.0	3208	15	US-10-210-130-38	Sequence 38, Appli	
19	29	100.0	3252	15	US-10-210-130-36	Sequence 36, Appli	
20	29	100.0	3262	15	US-10-179-381-4	Sequence 4, Appli	
21	29	100.0	3268	15	US-10-179-381-2	Sequence 2, Appli	
22	29	100.0	23	122	10	US-09-764-891-3273	
23	29	100.0	24	27	14	US-10-125-567A-438	
24	29	100.0	25	27	9	US-09-756-338A-2	
25	29	100.0	26	27	645	10	US-09-977-639A-28
26	29	100.0	27	27	645	10	US-09-977-418-28
27	29	100.0	27	27	645	10	US-09-977-418-46
28	29	100.0	28	27	645	10	US-09-977-033A-28
29	29	100.0	29	27	645	10	US-09-977-033A-16
30	29	100.0	30	27	645	10	US-09-977-751C-28
31	29	100.0	31	27	645	10	US-09-977-751C-46
32	29	100.0	32	27	645	10	US-09-977-639A-28
33	29	100.0	33	27	645	10	US-09-977-639A-46
34	29	100.0	34	27	645	11	US-09-977-819B-28
35	29	100.0	35	27	645	11	US-09-977-819B-46
36	29	100.0	36	27	645	14	US-10-282-837-2
37	29	100.0	37	27	645	14	US-10-145-585-2
38	29	100.0	38	27	683	10	US-09-977-418-26
39	29	100.0	39	27	683	10	US-09-977-033A-26
40	29	100.0	40	27	683	10	US-09-977-639A-26
41	29	100.0	41	27	683	11	US-09-977-819B-26
42	29	100.0	42	27	683	11	US-09-977-819B-26
43	29	100.0	43	27	683	9	US-09-826-366-7
44	29	100.0	44	27	683	9	US-09-903-320-4
45	29	100.0	45	27	690	9	US-09-903-0885-49

ALIGNMENTS

```
RESULT 1
US-09-848-834A-6
; Sequence 6, Application US/09848834A.

GENERAL INFORMATION:
; PATENT NO.: US2002007641A1
; APPLICANT: Aphton Corporation
; TITLE OF INVENTION: ChimERIC PeptidE Immunogens
; FILE REFERENCE: 1102865-0047
; CURRENT APPLICATION NUMBER: US/09/848,834A
; CURRENT FILING DATE: 2001-05-04
; PRIOR APPLICATION NUMBER: 60/202,328
; PRIOR FILING DATE: 2000-05-05
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: Patentin version 3.0
SEQ ID NO. 6
; LENGTH: 6
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic peptide
US-09-848-834A-6
```

FILE REFERENCE: 1102865-0047
 CURRENT APPLICATION NUMBER: US/09/848, 834A
 CURRENT FILING DATE: 2001-05-04
 PRIOR APPLICATION NUMBER: 60/202, 328
 PRIORITY FILING DATE: 2000-05-05
 NUMBER OF SEQ ID NOS: 20
 SEQ ID NO: 7
 LENGTH: 8
 TYPE: PRT
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: Synthetic peptide
 US-09-848-834A-7

Query Match Score 29; DB 9; Length 8;
 Best Local Similarity 100.0%; Pred. No. 7.1e+05;
 Matches 6; Conservative 0; Mismatches 0;
 Indels 0; Gaps 0;

Qy	1 SSGPSL 6
Db	1 SSGPSL 6

RESULT 3
 US-09-848-834A-15
 Sequence 15, Application US/09848834A
 Patent No. US2002007616A1
 GENERAL INFORMATION:
 APPLICANT: Abhton Corporation
 TITLE OF INVENTION: Chimeric Peptide Immunogens

FILE REFERENCE: 1102865-0047
 CURRENT APPLICATION NUMBER: US/09/848, 834A
 CURRENT FILING DATE: 2001-05-04
 PRIOR APPLICATION NUMBER: 60/202, 328
 PRIOR FILING DATE: 2000-05-05
 NUMBER OF SEQ ID NOS: 20
 SEQ ID NO: 15
 LENGTH: 31
 TYPE: PRT
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: Chimeric Peptide consisting of amino acid sequence 1-10 of the GnRH hormone linked by a spacer to amino acid sequence 830-844 of the Tetanus toxoid precursor (Tentoxilysin)

Qy	1 SSGPSL 6
Db	1 SSGPSL 6

Query Match Score 29; DB 9; Length 31;
 Best Local Similarity 100.0%; Pred. No. 7.7e+05;
 Matches 6; Conservative 0; Mismatches 0;
 Indels 0; Gaps 0;

RESULT 4
 US-09-848-834A-13
 Sequence 13, Application US/09848834A

Query Match Score 29; DB 9; Length 31;
 Best Local Similarity 100.0%; Pred. No. 7.7e+05;
 Matches 6; Conservative 0; Mismatches 0;
 Indels 0; Gaps 0;

RESULT 5
 US-09-848-834A-16
 Sequence 16, Application US/09848834A
 Patent No. US2002007616A1
 GENERAL INFORMATION:
 APPLICANT: Abhton Corporation
 TITLE OF INVENTION: Chimeric Peptide Immunogen
 FILE REFERENCE: 1102865-0047
 CURRENT APPLICATION NUMBER: US/09/848, 834A
 CURRENT FILING DATE: 2001-05-04
 PRIOR APPLICATION NUMBER: 60/202, 328
 PRIOR FILING DATE: 2000-05-05
 NUMBER OF SEQ ID NOS: 20
 SEQ ID NO: 16
 LENGTH: 36
 TYPE: PRT
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: Chimeric Peptide consisting of amino acid sequence 1-10 of the GnRH hormone linked by a spacer to amino acid sequence 378-398 of the Plasmodium falciparum circumsporozoite (CSP) protein.

Qy	1 SSGPSL 6
Db	1 SSGPSL 16

OTHER INFORMATION: Spacer peptide
 NAME/KEY: PEPTIDE (36)
 LOCATION: (17) . . .
 OTHER INFORMATION: Amino acid sequence 378-398 of the Malaria
 OTHER INFORMATION: (Plasmodium falciparum) circumsporozoite
 OTHER INFORMATION: (CSP) protein
 US-09-848-834A-16

Query Match Score 29; DB 9; Length 36;
 Best Local Similarity 100.0%; Pred. No. 89;
 Matches 6; Conservative 0; Mismatches 0;
 Indels 0; Gaps 0;

Qy 1 SSGPSL 6
 Db 11 SSGPSL 16

RESULT 6
 US-09-848-834A-14
 Sequence 14, Application US/09848834A.
 Patent No. US20020076416A1.

GENERAL INFORMATION:
 APPLICANT: Aphton Corporation
 TITLE OF INVENTION: Chimeric Peptide Immunogens
 FILE REFERENCE: 1102865_0047

CURRENT APPLICATION NUMBER: US/09/848,834A
 CURRENT FILING DATE: 2001-05-04
 PRIOR APPLICATION NUMBER: 6/202,328
 PRIOR FILING DATE: 2000-05-05
 SOFTWARE: PatentIn version 3.0
 SEQ ID NO 14
 LENGTH: 37
 TYPE: PRT
 ORGANISM: Artificial Sequence

FEATURE:
 OTHER INFORMATION: Chimeric Peptide consisting of amino acid sequence 1-10 of the
 OTHER INFORMATION: GnRH linked by a spacer to amino acid sequence 947-967 of
 OTHER INFORMATION: the Tetanus Toxoid precursor (TetoxoLysin)
 NAME/KEY: MOD RES
 LOCATION: (1) . . .(1)
 OTHER INFORMATION: Pyroglutamic acid or 5-oxopropine
 NAME/KEY: PEPTIDE
 LOCATION: (1) . . .(10)
 OTHER INFORMATION: Amino acid sequence 1-10 of the human GnRH hormone
 NAME/KEY: PEPTIDE
 LOCATION: (11) . . .(16)
 OTHER INFORMATION: Spacer peptide
 NAME/KEY: PEPTIDE
 LOCATION: (17) . . .(31)
 OTHER INFORMATION: Amino acid sequence 830-844 of the Tetanus toxoid precursor
 NAME/KEY: PEPTIDE
 LOCATION: (32) . . .(37)
 OTHER INFORMATION: Spacer peptide
 NAME/KEY: PEPTIDE
 LOCATION: (38) . . .(46)
 OTHER INFORMATION: Amino acid sequence 2-10 of the human GnRH hormone

US-09-848-834A-19

Query Match Score 100.0%; DB 9; Length 46;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 SSGPSL 6
 Db 11 SSGPSL 16

RESULT 8
 US-09-848-834A-17
 Sequence 17, Application US/09848834A
 Patent No. US20020076416A1.

GENERAL INFORMATION:
 APPLICANT: Aphton Corporation
 TITLE OF INVENTION: Chimeric Peptide Immunogens
 FILE REFERENCE: 1102865_0047

CURRENT APPLICATION NUMBER: US/09/848,834A
 CURRENT FILING DATE: 2001-05-04
 PRIOR APPLICATION NUMBER: 6/202,328
 PRIOR FILING DATE: 2000-05-05
 NUMBER OF SEQ ID NOS: 20
 SOFTWARE: PatentIn version 3.0
 SEQ ID NO 17
 LENGTH: 47
 TYPE: PRT
 ORGANISM: Artificial Sequence

FEATURE:
 OTHER INFORMATION: Chimeric Peptide consisting of amino acid sequence 1-10 of the
 OTHER INFORMATION: RH hormone linked by a spacer to amino acid sequence 288-302 of
 OTHER INFORMATION: the Measles virus protein F linked by a spacer to amino acid 5E
 OTHER INFORMATION: sequence 2-10 of the GnRH hormone
 NAME/KEY: MOD RES
 LOCATION: (1) . . .(1)
 OTHER INFORMATION: Pyroglutamic acid or 5-oxopropine
 NAME/KEY: MOD_RES

RESULT 7
 US-09-848-834A-19
 Sequence 19, Application US/09848834A
 Patent No. US20020076416A1.

GENERAL INFORMATION:
 APPLICANT: Aphton Corporation
 TITLE OF INVENTION: Chimeric Peptide Immunogens
 FILE REFERENCE: 1102865_0047

CURRENT APPLICATION NUMBER: US/09/848,834A
 CURRENT FILING DATE: 2001-05-04
 PRIOR APPLICATION NUMBER: 6/202,328
 PRIOR FILING DATE: 2000-05-05

APPLICANT: Hanzel, David K.
 APPLICANT: Chen, Wenheng
 TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
 TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY
 FILE REFERENCE: Aeonimica-X-1
 CURRENT APPLICATION NUMBER: US/09/864,761
 CURRENT FILING DATE: 2001-05-23
 PRIOR APPLICATION NUMBER: US 60/18C,312
 PRIOR FILING DATE: 2000-02-04
 PRIOR APPLICATION NUMBER: US 60/207,456
 PRIOR FILING DATE: 2000-05-26
 PRIOR APPLICATION NUMBER: US 09/632,366
 PRIOR FILING DATE: 2000-08-03
 PRIOR APPLICATION NUMBER: GB 24263,6
 PRIOR FILING DATE: 2000-10-04
 PRIOR APPLICATION NUMBER: US 60/236,359
 PRIOR FILING DATE: 2000-09-27
 PRIOR APPLICATION NUMBER: PCT/US01/00666
 PRIOR FILING DATE: 2001-01-10
 PRIOR APPLICATION NUMBER: PCT/US01/00667
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00664
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00669
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00665
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00668
 PRIOR FILING DATE: 2001-01-10
 PRIOR APPLICATION NUMBER: PCT/US01/00663
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00662
 PRIOR APPLICATION NUMBER: PCT/US01/00661
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00670
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: US 60/234,687
 PRIOR FILING DATE: 2000-09-1
 PRIOR FILING DATE: 2000-06-09/608,408
 PRIOR FILING DATE: 2000-06-09/774,203
 PRIOR FILING DATE: 2001-01-19
 NUMBER OF SEQ ID NOS: 49117
 SOFTWARE: Amomax Sequence Listing Engine vers. 1.1
 SEQ ID NO: 47649
 LENGTH: 53
 TYPE: PRT
 ORGANISM: Homo sapiens
 FEATURE:
 OTHER INFORMATION: MAP TO 283826_12
 OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 0.73
 OTHER INFORMATION: EXPRESSED IN HBL010, SIGNAL = 0.96
 OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 0.57
 OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 0.62
 OTHER INFORMATION: SWISSPROT HIT: OS399, EVALUATE 5.90e+00
 OTHER INFORMATION: EST_HUMAN HIT: AAC13575_1, EVALUATE 3.00e-22
 US-09-864-761-47649

Query Match 100.0%; Score 29; DB 9; Length 53;
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 12
 US-09-731-872-425
 i Sequence 425, APP10, Application US/09731872
 i Patent No. US2002012604x1
 i GENERAL INFORMATION:

TITLE OF INVENTION: LUMAZINE SYNTHASE AND
 RIBOFLAVIN SYNTHASE

APPLICANT: Dumas Milne Edwards, Jean Baptiste
 APPLICANT: Bougueret, Lydie
 APPLICANT: Jober, Severin
 TITLE OF INVENTION: FULL-LENGTH HUMAN cDNAs ENCODING POTENTIALLY SECRETED PROTEINS
 FILE REFERENCE: 79, US3, REG
 CURRENT APPLICATION NUMBER: US/09/731,872
 CURRENT FILING DATE: 2000-12-07
 PRIOR APPLICATION NUMBER: US 60/169,629
 PRIOR FILING DATE: 1999-12-08
 PRIOR APPLICATION NUMBER: US 60/187,470
 PRIOR FILING DATE: 2000-03-06
 NUMBER OF SEQ ID NOS: 482
 SOFTWARE: Patent.pn
 SEQ ID NO: 425
 LENGTH: 75
 TYPE: PRT
 ORGANISM: Homo sapiens
 US-09-731-872-425

Query Match 100.0%; Score 29; DB 9; Length 75;
 Best Local Similarity 100.0%; Pred. No. 1.8e+02;
 Matches 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 13
 US-09-776-997-425
 Sequence 425, Application US/09876997
 Publication No. US2003015221A1
 GENERAL INFORMATION:
 APPLICANT: Dumas Milne Edwards, Jean Baptiste
 APPLICANT: Bougueret, Lydie
 APPLICANT: Jober, Severin
 TITLE OF INVENTION: FULL-LENGTH HUMAN cDNAs ENCODING POTENTIALLY SECRETED PROTEINS
 FILE REFERENCE: 78, US4, CIP
 CURRENT APPLICATION NUMBER: US/09/876,997
 CURRENT FILING DATE: 2001-06-08
 PRIOR APPLICATION NUMBER: US 09/731,872
 PRIOR FILING DATE: 2000-12-07
 PRIOR APPLICATION NUMBER: US 60/187,470
 PRIOR FILING DATE: 2000-03-06
 PRIOR APPLICATION NUMBER: US 60/169,629
 PRIOR FILING DATE: 1999-12-08
 NUMBER OF SEQ ID NOS: 482
 SOFTWARE: Patent.pn
 SEQ ID NO: 425
 LENGTH: 75
 TYPE: PRT
 ORGANISM: Homo sapiens
 US-09-876-997-425

Query Match 100.0%; Score 29; DB 10; Length 75;
 Best Local Similarity 100.0%; Pred. No. 1.8e+02;
 Matches 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 14
 US-09-870-756-38
 Sequence 38, Application US/09870756
 i Patent No. US20020052023A1
 i GENERAL INFORMATION:
 i APPLICANT: Viitanen, Paul Veikko
 i BACOT, Karen Onley
 i JORDAN, Douglas Brian
 i TITLE OF INVENTION: LUMAZINE SYNTHASE AND
 RIBOFLAVIN SYNTHASE

NUMBER OF SEQUENCES: 39
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: E. I. DU PONT DE NEMOURS AND COMPANY
 STREET: 1007 MARKET STREET
 CITY: WILMINGTON
 STATE: DELAWARE
 COUNTRY: UNITED STATES OF AMERICA
 ZIP: 19898
 COMPUTER READABLE FORM:
 MEDIUM TYPE: DISKETTE, 3.50 INCH
 COMPUTER: IBM PC COMPATIBLE
 OPERATING SYSTEM: MICROSOFT WORD FOR WINDOWS 95
 SOFTWARE: MICROSOFT WORD VERSION 7.0A
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/09/870,756
 FILING DATE: 31-May-2001
 ATTORNEY/AGENT INFORMATION:
 NAME: FLOYD LINDA AXAMETHY
 REGISTRATION NUMBER: 33,692
 REFERENCE/DOCKET NUMBER: CL-1083
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 302-992-8112
 TELEFAX: 302-773-0164
 INFORMATION FOR SEQ ID NO: 38:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 200 amino acids
 TYPE: amino acid
 STRANDEDNESS: not relevant
 TOPOLOGY: not relevant
 MOLECULE TYPE: protein
 ORIGINAL SOURCE:
 INDIVIDUAL ISOLATE: M. grisea LS
 SEQUENCE DESCRIPTION: SEQ ID NO: 38:
 US-09-870-756-38

Query Match Score 29; DB 9; Length 200;
 Best Local Similarity 100.0%; Pred. No. 4.5e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 SSGPSL 6
 Db 79 SSGPSL 84

RESULT 15
 US-09-874-585B-38
 Sequence 38, Application US/09874585B
 Patient No. US20020127670A1
 GENERAL INFORMATION:
 APPLICANT: Viitanen, Paul Veikko
 APPLICANT: Jordan, Douglas Brian
 APPLICANT: Bacot, Karen Onley
 TITLE OF INVENTION: Riboflavin Synthase Genes and Enzymes and Methods of Use
 FILE REFERENCE: CL1083 US DIV2
 CURRENT APPLICATION NUMBER: US/09/874,585B
 CURRENT FILING DATE: 2002-03-05
 PRIOR APPLICATION NUMBER: 09/874,585
 PRIOR FILING DATE: 2001-06-05
 PRIOR APPLICATION NUMBER: 08/912,218
 PRIOR FILING DATE: 1997-08-15
 SOFTWARE: Microsoft Office 97
 SEQ ID NO 38
 LENGTH: 200
 TYPE: FRT
 ORGANISM: Magnaporthe grisea
 US-09-874-585B-38

Query Match Score 29; DB 9; Length 200;
 Best Local Similarity 100.0%; Pred. No. 4.5e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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OM protein - protein search, using sw model

Run on: March 10, 2004, 08:58:48 ; Search time 9.03502 Seconds
 (without alignments)
 187.635 Million cell updates/sec

Title: US-09-848-834A-6

Perfect score: 29

Sequence: 1 SSGPSL 6

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 15861.07 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 15861.07

Minimum DB seq length: 0
 Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
 Maximum Match 100%
 Listing first 45 summaries

Database : A_Geneseq_29Jan04:
 1: GeneseqP1900s:
 2: GeneseqP1900s:
 3: GeneseqP2000s:
 4: GeneseqP2001s:
 5: GeneseqP2002s:
 6: GeneseqP2003as:
 7: GeneseqP2003bs:
 8: GeneseqP2004s:
 Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query	Score	Match	Length	DB ID	Description
1	AAU11417	6	5	6	AAU11417	Aau11417 Synthetic
2	AATU1418	5	5	6	AAU11418	Aau11418 Synthetic
3	AATU1426	31	5	31	AAU11426	Aau11426 Synthetic
4	AATU1424	34	5	34	AAU11424	Aau11424 Synthetic
5	AATU1427	36	5	36	AAU11427	Aau11427 Synthetic
6	AAU11425	37	5	37	AAU11425	Aau11425 Synthetic
7	AATU1430	46	5	46	AAU11430	Aau11430 Synthetic
8	AATU1428	47	5	47	AAU11428	Aau11428 Synthetic
9	AATU1429	50	5	50	AAU11429	Aau11429 Synthetic
10	AAG03447	51	3	51	AAG03447	Aag03447 Human sec
11	AATU1431	51	5	51	AAU11431	Aau11431 Synthetic
12	AAM1164	53	4	53	AAM1164	Abb1164 Peptide #
13	ABB37195	53	4	53	ABB37195	Abb37195 Peptide #
14	AAM70325	53	4	53	AAM70325	Aam70325 Human bon
15	AAM05786	53	4	53	AAM05786	Aam05786 Peptide #
16	AAG89305	75	4	75	AAG89305	Aag89305 Human sec
17	AAQ10754	117	4	117	AAQ10754	Aaq10754 Human pol
18	AAV73871	128	2	128	AAV73871	Aav73871 Human Pro
19	AATU14353	100	0	100	AATU14353	Aau42353 Propionib
20	ABM38872	172	4	172	ABM38872	Abm38872 Propionib
21	AAB30732	100	0	100	AAB30732	Aab30732 Novel hum
22	ABG19555	181	4	181	ABG19555	Abg19555 Novel hum
23	ABG19556	100	0	100	ABG19556	Abg19556 Novel hum
24	AAV7393	187	4	187	AAV7393	Aav7393 Magnapor
25	AAB655772	100	0	100	AAB655772	Aab655772 Drosophil

ALIGNMENTS

RESULT 1

AAU11417 standard; peptide; 6 AA.

ID AAU11417

XX AC AAU11417:

XX DT 12-MAR-2002 (first entry)

XX DE Synthetic spacer peptide #2.

XX KW Gonadotrophin releasing hormone; GnRH; synthetic immunogen;

XX KW luteinising hormone releasing hormone; LHRH; contraceptive;

XX KW promising helper T-cell peptide epitope; immunomimetic peptide epitope;

XX KW breast cancer; uterine cancer; gynaecological cancer; endometriosis;

XX KW uterine fibroid; benign prostatic hypertrophy; prostate cancer;

XX KW spacer peptide.

OS Synthetic.

XX PN WO200185763-A2.

XX PD 15-NOV-2001.

XX PP 04-MAY-2001; 2001WO-US014463.

XX PR 05-MAY-2000; 2000US-0202328P.

XX PA (APHT-) APHTON CORP.

XX PI Grimes S, Michaelis D, Stevens VC;

XX DR WPI; 2002-049440/06.

XX PT Novel synthetic immunogen for inducing immune response against

PT gonadotropin releasing hormone, comprises fusion peptide having

PT promiscuous helper T-cell peptide epitope and immunomimetic peptide epitope

PT or its analog.

XX PS Claim 10; Page 6; 43pp; English.

XX CC The invention relates to a synthetic immunogen for inducing specific

CC antibodies against gonadotropin releasing hormone (GnRH) also known as

CC luteinising hormone releasing hormone (LHRH) comprising a fusion peptide

CC which comprises a promiscuous helper T-cell peptide epitope and

CC immunomimetic peptide epitope or its analogue. The synthetic immunogen is

CC useful inducing an immune response against GnRH in an animal subject, and

CC such is useful as a contraceptive and in the treatment of diseases

CC such as cancer (of the breast, uterus and other gynaecological cancer),

CC

CC endometriosis, uterine fibroids, benign prostatic hypertrophy and
 CC prostate cancer. The immunogen is effective in eliciting high and
 CC specific anti-GnRH antibody titres. The present sequence is a synthetic
 CC spacer peptide used in the immunogen of the invention
 XX Sequence 6 AA;

Query Match	100.0%	Score 29;	DB 5;	Length 6;		Matches 6;	Conservative	0;	Mismatches 0;	Indels 0;	Gaps 0;
Best Local Similarity	100.0%	Pred. No.	1.4e+06;			Qy	1 SSGSLS 6				
Matches 6	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;		ID	AAU1426 standard; peptide; 31 AA.				
Qy	1 SSGSLS 6					Db	1 SSGSLS 6				
Db	1 SSGSLS 6										

RESULT 2
 AAU1418
 ID AAU1418 standard; peptide; 8 AA.
 XX
 AC AAU1418;
 XX DT 12-MAR-2002 (first entry)
 XX Synthetic spacer peptide #3.
 XX Gonadotrophin releasing hormone; GnRH; synthetic immunogen;
 DE luteinising hormone releasing hormone; LHRH; contraceptive;
 XX promiscuous helper T-cell peptide epitope; immunomimic Peptide epitope;
 KW breast cancer; uterine cancer; gynaecological cancer; endometriosis;
 KW uterine fibroid; benign prostatic hypertrophy; prostate cancer;
 KW spacer peptide.
 XX Synthetic.
 XX WO200185763-A2.
 PN 15-NOV-2001.
 XX PD 15-NOV-2001.
 XX PP 04-MAY-2001; 2001WO-US014363.
 XX PR 05-MAY-2000; 2000US-0202328P.
 PA (APHT-) APTHON CORP.
 PI Grimes S, Michaeli D, Stevens VC;
 XX DR WPI; 2002-049440/06.
 XX PS Claim 10; Page 6; 43PP; English.
 CC The invention relates to a synthetic immunogen for inducing specific
 CC antibodies against gonadotropin releasing hormone (GnRH also known as
 CC luteinising hormone releasing hormone, LHRH) comprising a fusion peptide
 CC which comprises a promiscuous helper T-cell peptide epitope and
 CC immunomimic peptide epitope or its analogue. The synthetic immunogen is
 CC useful in inducing an immune response against GnRH in an animal subject, and
 CC as such is useful as a contraceptive and in the treatment of diseases
 CC such as cancer (of the breast, uterus and other gynaecological cancer),
 CC endometriosis, uterine fibroids, benign prostatic hypertrophy and
 CC prostate cancer. The immunogen is effective in eliciting high and
 CC specific anti-GnRH antibody titres. The present sequence is a synthetic
 CC spacer peptide used in the immunogen of the invention
 XX Sequence 8 AA;

Query Match	100.0%	Score 29;	DB 5;	Length 8;		Matches 6;	Conservative	0;	Mismatches 0;	Indels 0;	Gaps 0;
Best Local Similarity	100.0%	Pred. No.	1.4e+06;			Qy	1 SSGSLS 6				
XX						ID	AAU1426				
						Db	1 SSGSLS 6				

RESULT 3
 AAU1426
 ID AAU1426 standard; peptide; 31 AA.
 XX
 AC AAU1426;
 XX DT 12-MAR-2002 (first entry)
 XX Synthetic immunogen peptide 7.
 XX Gonadotrophin releasing hormone; GnRH; synthetic immunogen;
 KW luteinising hormone releasing hormone; LHRH; contraceptive;
 KW promiscuous helper T-cell peptide epitope; immunomimic Peptide epitope;
 KW breast cancer; uterine cancer; gynaecological cancer; endometriosis;
 KW uterine fibroid; benign prostatic hypertrophy; prostate cancer.
 XX Clostridium tetani.
 OS Mammalia.
 OS Synthetic.
 OS Chimeric.
 XX Key
 FT Peptide 1..10 "Gonadotrophin releasing hormone epitope"
 FT Misc-difference 1 /label= OTHER
 FT Peptide /note= "Other= Pyro-glutamic acid or 5-oxo proline"
 FT Peptide 11..16
 FT Peptide /note= "Spacer Peptide"
 FT Peptide 17..31 "Tetanus toxoid sequence (830-844 aa)"
 FT Peptide /note= "Tetanus toxoid sequence (830-844 aa)"
 XX WO200185763-A2.
 XX PD 15-NOV-2001.
 XX PP 04-MAY-2001; 2001WO-US014363.
 XX PR 05-MAY-2000; 2000US-0202328P.
 XX PA (APHT-) APTHON CORP.
 XX PI Grimes S, Michaeli D, Stevens VC;
 XX DR WPI; 2002-049440/06.
 XX PS Claim 10; Page 10; 43PP; English.
 CC The invention relates to a synthetic immunogen for inducing specific
 CC antibodies against gonadotropin releasing hormone (GnRH also known as
 CC luteinising hormone releasing hormone, LHRH) comprising a fusion peptide
 CC which comprises a promiscuous helper T-cell peptide epitope and
 CC immunomimic peptide epitope or its analogue. The synthetic immunogen is
 CC useful in inducing an immune response against GnRH in an animal subject, and
 CC as such is useful as a contraceptive and in the treatment of diseases
 CC such as cancer (of the breast, uterus and other gynaecological cancer),
 CC endometriosis, uterine fibroids, benign prostatic hypertrophy and
 CC prostate cancer. The immunogen is effective in eliciting high and
 CC specific anti-GnRH antibody titres. The present sequence is a synthetic
 CC spacer peptide used in the immunogen of the invention
 XX Sequence 8 AA;

Query Match	100.0%	Score 29;	DB 5;	Length 8;		Matches 6;	Conservative	0;	Mismatches 0;	Indels 0;	Gaps 0;
Best Local Similarity	100.0%	Pred. No.	1.4e+06;			Qy	1 SSGSLS 6				
XX						ID	AAU1426				
						Db	1 SSGSLS 6				

Claim 11; Page 10; 43PP; English.

The invention relates to a synthetic immunogen for inducing specific antibodies against gonadotropin releasing hormone (GnRH also known as luteinising hormone releasing hormone, LHRH) comprising a fusion peptide which comprises a promiscuous helper T-cell peptide epitope and immunomimic peptide epitope or its analogue. The synthetic immunogen is useful in inducing an immune response against GnRH in an animal subject, and as such is useful as a contraceptive and in the treatment of diseases such as cancer (of the breast, uterus and other gynaecological cancer), endometriosis, uterine fibroids, benign prostatic hypertrophy and prostate cancer. The immunogen is effective in eliciting high and specific anti-GnRH antibody titres. The present sequence is a synthetic immunogen of the invention

PT Grimes S, Michaeli D, Stevens VC;
 XX WPI; 2002-049440/06.

XX Novel synthetic immunogen for inducing immune response against
 PT gonadotropin releasing hormone, comprises fusion peptide having
 PT promiscuous helper T-cell peptide epitope and immunomimic peptide epitope
 or its analog.
 XX
 ES Claim 11; Page 12; 43PP; English.

XX The invention relates to a synthetic immunogen for inducing specific
 CC antibodies against gonadotropin releasing hormone (GnRH also known as
 CC luteinising hormone releasing hormone, LHRH) comprising a fusion peptide
 CC which comprises a promiscuous helper T-cell peptide epitope and
 CC immunomimic peptide epitope or its analogue. The synthetic immunogen is
 CC useful inducing an immune response against GnRH in an animal subject, and
 CC as such is useful as a contraceptive and in the treatment of diseases
 CC such as cancer (of the breast, uterus and other gynaecological cancer),
 CC endometriosis, uterine fibroids, benign prostatic hypertrophy and
 CC prostate cancer. The immunogen is effective in eliciting high and
 CC specific anti-GnRH antibody titres. The present sequence is a synthetic
 CC immunogen of the invention
 XX

SQ Sequence 46 AA:
 Query Match Score 29; DB 5; Length 46;
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 SSSPSL 6
 Db 11 SSSPSL 16

RESULT 8
 ID AAU11428 standard; peptide; 47 AA.
 XX AAU11428;
 AC AAU11428;
 XX DT 12-MAR-2002 (first entry)
 XX DE Synthetic immunogen peptide 9.
 XX Gonadotrophin releasing hormone; GnRH; synthetic immunogen;
 KW luteinising hormone releasing hormone; LHRH; contraceptive;
 KW promiscuous helper T-cell peptide epitope; immunomimic peptide epitope;
 KW breast cancer; uterine cancer; gynaecological cancer; endometriosis;
 KW uterine fibroid; benign prostatic hypertrophy; prostate cancer.
 OS Plasmodium falciparum.
 OS Mammalia.
 OS Chimeric.
 XX Key Location/Qualifiers
 PT Peptide 1..10 /note= "Gonadotrophin releasing hormone epitope (1..10 aa)"
 PT Misc-difference 1 /label= OTHER /note= "Other= Pyro-glutamic acid or 5-oxo proline"
 PT Peptide 11..16 /note= "Spacer peptide"
 PT Peptide 17..34 /note= "Malaria CSP protein (288-302 aa)"
 PT Peptide 35..39 /note= "Spacer peptide"
 PT Peptide 39..47 /note= "Gonadotrophin releasing hormone epitope (2-10 aa)"
 PT Modified-site 47

FT /note= "Amidated glycine or glycynamide"
 XX
 DR WO200185763-A2.
 XX PD 15-NOV-2001.
 XX PF 04-MAY-2001; 2001WO-US014363.
 XX PR 05-MAY-2000; 2000US-0202328P.
 XX (APHT-) APHTON CORP.
 XX PA
 PI Grimes S, Michaeli D, Stevens VC;
 XX DR 2002-049440/06.
 XX PT Novel synthetic immunogen for inducing immune response against
 PT gonadotropin releasing hormone, comprises fusion peptide having
 PT promiscuous helper T-cell peptide epitope and immunomimic peptide epitope
 PT or its analog.
 XX Claim 11; Page 11; 43PP; English.
 XX The invention relates to a synthetic immunogen for inducing specific
 CC antibodies against gonadotropin releasing hormone (GnRH also known as
 CC luteinising hormone releasing hormone, LHRH) comprising a fusion peptide
 CC which comprises a promiscuous helper T-cell peptide epitope and
 CC immunomimic peptide epitope or its analogue. The synthetic immunogen is
 CC useful inducing an immune response against GnRH in an animal subject, and
 CC as such is useful as a contraceptive and in the treatment of diseases
 CC such as cancer (of the breast, uterus and other gynaecological cancer),
 CC endometriosis, uterine fibroids, benign prostatic hypertrophy and
 CC prostate cancer. The immunogen is effective in eliciting high and
 CC specific anti-GnRH antibody titres. The present sequence is a synthetic
 CC immunogen of the invention
 XX

SQ Sequence 47 AA:
 Query Match Score 29; DB 5; Length 47;
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 SSGRSL 6
 Db 11 SSGRSL 16

RESULT 9
 AAU11429 standard; peptide; 50 AA.
 XX AAU11429;
 AC AAU11429;
 XX DT 12-MAR-2002 (first entry)
 XX DE Synthetic immunogen peptide 10.
 XX KW Gonadotrophin releasing hormone; GnRH; synthetic immunogen;
 KW luteinising hormone releasing hormone; LHRH; contraceptive;
 KW promiscuous helper T-cell peptide epitope; immunomimic peptide epitope;
 KW breast cancer; uterine cancer; gynaecological cancer; endometriosis;
 KW uterine fibroid; benign prostatic hypertrophy; prostate cancer.
 OS Clostridium tetani.
 OS Mammalia.
 OS Synthetic.
 OS Chimeric.
 XX Key Location/Qualifiers
 PT Peptide 1..10 /note= "Gonadotrophin releasing hormone epitope (1..10 aa)"
 PT Peptide 11..16 /note= "Spacer peptide"
 PT Peptide 17..34 /note= "Malaria CSP protein (288-302 aa)"
 PT Peptide 35..39 /note= "Spacer peptide"
 PT Peptide 39..47 /note= "Gonadotrophin releasing hormone epitope (1..10 aa)"
 PT Misc-difference 1

FT /label= OTHER
 FT /note= "Other= Pyro-glutamic acid or 5-oxo proline"
 FT 11. .16
 FT /note= "Spacer peptide"
 FT 17. .37
 FT Peptide /note= "Tetanus toxoid (947-967 aa)"
 FT 38. .41
 FT /note= "Spacer peptide"
 FT 42. .50
 FT /note= "Gonadotrophin releasing hormone epitope (2-10 aa)"
 FT 50
 FT Modified-site /note= "Amidated glycine or glycynamide"
 FT WO200185763-A2.
 XX 15-NOV-2001.
 PD 04-MAY-2001; 2001WO-US014333.
 PR 05-MAY-2000; 2000US-0202328P.
 XX (APHT-) APHTON CORP.
 PA Grimes S, Michaeli D, Stevens VC;
 PI XX 2002-049440/06.
 DR Novel synthetic immunogen for inducing immune response against
 PR gonadotropin releasing hormone, comprises fusion peptide having
 XX promiscuous helper T-cell peptide epitope and immunomimic peptide epitope
 PR or its analog.
 XX Claim 11; Page 11; 43PP; English.
 PS The invention relates to a synthetic immunogen for inducing specific
 CC antibodies against gonadotropin releasing hormone (GnRH) also known as
 CC luteinising hormone releasing hormone (LHRH) comprising a fusion peptide
 CC which comprises a promiscuous helper T-cell peptide epitope and
 CC immunomimic peptide epitope or its analogue. The synthetic immunogen is
 CC useful inducing an immune response against GnRH in an animal subject, and
 CC as such is useful as a contraceptive and in the treatment of diseases
 CC such as cancer (of the breast, uterus and other gynaecological cancer),
 CC endometriosis, uterine fibroids, benign prostatic hyper trophy and
 CC prostate cancer. The immunogen is effective in eliciting high and
 CC specific anti-GnRH antibody titres. The present sequence is a synthetic
 CC immunogen of the invention
 XX Sequence 50 AA;
 SQ Query Match 100.0%; Score 29; DB 5; Length 50;
 Best Local Similarity 100.0%; Pred. No. 1.4e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 AC QY 1 SSGPSL 6
 AC DB 11 SSGPSL 16
 AC DT 06-OCT-2000 (first entry)
 AC Human secreted protein, SEQ ID NO: 7528.
 AC XX Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;
 AC XX Gene therapy; chromosome mapping.
 AC XX Homo sapiens.
 PN XX EP1033401-A2.
 PN XX 06-SEP-2000.
 PD XX 21-FEB-2000; 2000EP-00200610.
 PR XX 26-FEB-1999; 99US-0122487P.
 PA XX (GEST) GENSET.
 PI XX Dumas Milne Edwards J, Duclert A, Giordano J;
 XX WPI; 2000-500381/45.
 DR XX N-BSDB; AAC03453.
 PT New nucleic acid that is a 5' expressed sequence tag (5' EST) for
 PT obtaining cDNAs and genomic DNAs that correspond to 5' ESTs and for
 PT diagnostic, forensic, gene therapy and chromosome mapping procedures.
 PR XX Claim 13; SEQ ID NO 7528; 71PP + Sequence Listing; English.
 PS XX The present sequence is a polypeptide encoded by one of a large number of
 CC ESTs derived from mRNAs encoding secreted proteins. The 5' ESTs were
 CC prepared from total human RNAs or PolyA+ RNAs derived from 30 different
 CC tissues. EST sequences usually correspond mainly to the 3' untranslated
 CC region (UTR) of the mRNA because they are often obtained from oligo-dT
 CC printed cDNA libraries. Such ESTs are not well suited for isolating cDNA
 CC sequences derived from the 5' ends of mRNAs and even in those cases where
 CC longer cDNA sequences have been obtained, the full 5' UTR is rarely
 CC included. 5' ESTs are derived from mRNAs with intact 5' ends and can
 CC therefore be used to obtain full length cDNAs and genomic DNAs. 5' ESTs
 CC are also used in diagnostic, forensic, gene therapy and chromosome
 CC mapping procedures. They are used to obtain upstream regulatory sequences
 CC and to design expression and secretion vectors
 SQ Sequence 51 AA;
 SQ Query Match 100.0%; Score 29; DB 3; Length 51;
 Best Local Similarity 100.0%; Pred. No. 1.4e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 AC QY 1 SSGPSL 6
 AC DB 16 SSGPSL 21
 AC DT 12-MAR-2002 (first entry)
 AC XX Synthetic immunogen peptide 12.
 AC XX AAU11431 standard; peptide; 51 AA.
 AC XX AAU11431;
 AC XX DT 12-MAR-2002
 AC XX KW Gonadotrophin releasing hormone; GnRH; synthetic immunogen;
 AC XX KW Luteinising hormone releasing hormone; LHRH; contraceptive;
 AC XX KW Promiscuous helper T-cell peptide epitope; immunomimic peptide epitope;
 AC XX KW Breast cancer; uterine cancer; gynaecological cancer; endometriosis;
 AC XX KW Uterine fibroid; benign prostatic hypertrophy; prostate cancer.
 AC OS Plasmodium falciparum.
 AC OS Mammalia.
 AC OS Synthetic.
 AC OS Chimeric.
 AC XX Key Peptide
 AC XX Location/Qualifiers
 AC XX 1..10
 AC XX /note= "Gonadotrophin releasing hormone epitope (1..10 aa)"
 AC XX Misc-difference 1
 AC XX FT

PT /label= OTHER
 PT /note= "Other= Pyro-glutamic acid or 5-oxo proline"
 PT 11 . 16
 PT /note= "Spacer peptide"
 PT 17 . 36
 PT /note= "Malaria CSP protein (378-398 aa)"
 PT 37 . 42
 PT /note= "Spacer Peptide"
 PT 43 . 51
 PT /note= "Gonadotrophin releasing hormone epitope (2-10 aa)"
 PT 51
 PT Modified-site /note= "Amidated glycine or glycynamide"
 PT WO200185763-A2.
 PN XX
 PD 15-NOV-2001.
 XX
 XX
 PR 04-MAY-2001; 2001WO-US014363.
 XX
 PR 05-MAY-2000; 2000US-0202328P.
 XX
 PA (APHT-) APHTON CORP.
 XX
 PI Grimes S, Michaeli D, Stevens VC;
 XX
 DR WPI; 2002-049440/06.
 XX
 PT Novel synthetic immunogen for inducing immune response against gonadotropin releasing hormone, comprises fusion peptide having promiscuous helper T-cell peptide epitope and immunomimic peptide epitope or its analog.
 XX
 Claim 11; Page 12-13; 43pp; English.
 CC The invention relates to a synthetic immunogen for inducing specific antibodies against Gonadotropin releasing hormone (GnRH) also known as luteinising hormone releasing hormone, LHRH comprising a fusion peptide which comprises a promiscuous helper T-cell peptide epitope and immunomimic peptide epitope or its analogue. The synthetic immunogen is useful inducing an immune response against GnRH in an animal subject, and as such is useful as a contraceptive and in the treatment of diseases, such as cancer (of the breast, uterus and other gynaecological cancer), endometriosis, uterine fibroids, benign prostatic hyper trophy and prostate cancer. The immunogen is effective in eliciting high and specific anti-GnRH antibody titres. The present sequence is a synthetic immunogen of the invention
 XX
 SQ Sequence 51 AA;

Query Match 100.0%; Score 29; DB 5; Length 51;
 Best Local Similarity 100.0%; Pred. No. 1.4e+05;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 SSGPSL 6
 Db 11 SSGPSL 16

RESULT 12
 ID AAM18164 standard; protein; 53 AA.
 XX
 AC AAM18164;
 XX
 DT 12-OCT-2001 (first entry)
 XX
 RESULT 12
 ID AAM18164 standard; protein; 53 AA.
 XX
 AC AAM18164;
 XX
 DT 12-OCT-2001 (first entry)
 XX
 DE Peptide #4598 encoded by probe for measuring cervical gene expression.
 XX
 Probe; human; microarray; gene expression; cervical epithelial cell;
 XX
 cervical cancer.
 OS Homo sapiens.

XX WO200157278-A2.
 PN XX
 PD 09-AUG-2001.
 XX
 PR 30-JAN-2001; 2001WO-US000670.
 XX
 PR 04-FEB-2000; 2000US-0180312P.
 PR 26-MAY-2000; 2000US-0207456P.
 PR 30-JUN-2000; 2000US-00623668.
 PR 03-AUG-2000; 2000US-00623666.
 PR 21-SEP-2000; 2000US-0234687P.
 PR 27-SEP-2000; 2000US-02363359P.
 PR 04-OCT-2000; 2000GB-00024263.
 XX
 PA (MOLEC-) MOLECULAR DYNAMICS INC.
 XX
 PI Penn SG, Hanelz DK, Chen W, Rank DR;
 XX
 DR WPI; 2001-488901/53.
 XX
 PT Human genome-derived single exon nucleic acid probes useful for analyzing gene expression in human cervical epithelial cells.
 XX
 PT
 XX
 PS Claim 27; SEQ ID NC 229900; 487pp; English.
 XX
 CC The present invention relates to human single exon nucleic acid probes (SNPs; see AA10066-AA128459). The present sequence is a peptide encoded by one such probe. The SNPs are derived from human HeLa cells. The SNPs can be used to produce a single exon microarray, which can be used for measuring human gene expression in a sample derived from human cervical epithelial cells. By measuring gene expression, the probes are therefore useful in grading and/or staging of diseases of the cervix, notably cervical cancer. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/pct sequences

XX
 SQ Sequence 53 AA;

Query Match 100.0%; Score 29; DB 4; Length 53;
 Best Local Similarity 100.0%; Pred. No. 1.5e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 SSGPSL 6
 Db 5 SSGPSL 10

RESULT 13
 ABB37195
 ID ABB37195 standard; peptide; 53 AA.
 XX
 AC ABB37195;
 XX
 DT 04-FEB-2002 (first entry)
 XX
 DE Peptide #4701 encoded by human foetal liver single exon probe.
 XX
 KW Human; foetal liver; gene expression; single exon nucleic acid probe.
 XX
 OS Homo sapiens.
 XX
 PN WO200157277-A2.
 XX
 PD 09-AUG-2001.
 XX
 PR 30-JAN-2001; 2001WO-US000669.
 XX
 PR 04-FEB-2000; 2000US-0180312P.
 PR 26-MAY-2000; 2000US-0207456P.
 PR 30-JUN-2000; 2000US-00623668.
 PR 03-AUG-2000; 2000US-00623666.
 PR 21-SEP-2000; 2000US-0234687P.

XX The present invention provides a number of single exon nucleic acid
 CC probes which are derived from genomic sequences expressed in the human
 CC bone marrow. They can be used to measure gene expression in bone marrow
 CC samples, which may enable the improved diagnosis and treatment of cancers
 CC such as lymphoma, leukaemia and myeloma. The present sequence is a
 CC protein encoded by one of the probes of the invention
 XX
 XX SQ Sequence 53 AA:
 Query Match 100.0%; Score 29; DB 4; Length 53;
 Best Local Similarity 100.0%; Pred. No. 1.5e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 YY 1 SSGPSSL 6
 Db 5 SSGPSSL 10
 RESULT 15
 AAM05786 standard; protein; 53 AA.
 ID AAM05786
 XX AC AAM05786;
 XX DT 09-OCT-2001 (First entry)
 XX DE Peptide #4468 encoded by probe for measuring breast gene expression.
 XX KW Probe; human; breast disease; breast cancer; development disorder;
 XX inflammatory disease; proliferative breast disease; non-carcinoma tumour.
 XX OS Homo sapiens.
 XX PN WO200157270-A2.
 XX PD 09-AUG-2001.
 XX XX 29-JAN-2001; 2001WO-US000661.
 XX PR 04-FEB-2000; 2000US-0180312P.
 XX PR 26-MAY-2000; 2000US-0207456P.
 XX PR 30-JUN-2000; 2000US-0060808.
 XX PR 03-AUG-2000; 2000US-00612166.
 XX PR 21-SEP-2000; 2000US-0234681P.
 XX PR 27-SEP-2000; 2000US-0236339P.
 XX PR 04-OCT-2000; 2000GB-00024263.
 XX PA (NOEB-) MOLECULAR DYNAMICS INC.
 XX PI Penn SG, Hanzel DK, Chen W, Rank DR;
 XX DR; 2001-476286/51.
 XX Novel single exon nucleic acid probe used to measuring gene expression in
 XX a human breast.
 XX Claim 27; SEQ ID NO 14526; 322DP; English.
 XX The present invention relates to novel single exon nucleic acid probes
 CC such probe. The probes are useful for measuring human gene expression in
 CC a human breast sample, where the probe hybridises at high stringency to a
 CC nucleic acid expressed in the human breast. The probes are useful for
 CC predicting, diagnosing, grading, staging, monitoring and prognosis
 CC diseases of the human breast, particularly those diseases with polygenic
 CC aetiology. The diseases include: breast cancer, disorders of development,
 CC inflammatory diseases of the breast, fibrotic changes, proliferative
 CC breast disease and non-carcinoma tumours. Note: The sequence data for
 CC this patent did not form part of the printed specification, but was
 CC obtained in electronic format directly from WIPO at
 CC ftp://wipo.int/pub/published_pct_sequences
 XX
 XX SQ Sequence 53 AA:
 Query Match 100.0%; Score 29; DB 4; Length 53;
 Best Local Similarity 100.0%; Pred. No. 1.5e+02;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 YY 1 SSGPSSL 6
 Db 5 SSGPSSL 10
 RESULT 14
 AAMM70325 standard; protein; 53 AA.
 ID AAMM70325;
 XX 06-NOV-2001 (first entry)
 XX Human bone marrow expressed probe encoded protein SEQ ID NO: 30631.
 XX Human; bone marrow expressed exon; gene expression analysis; probe;
 XX microarray; cancer; leukaemia; lymphoma; myeloma.
 XX Homo sapiens.
 XX WO200157276-A2.
 XX PD 09-AUG-2001.
 XX 30-JAN-2001; 2001WO-US000668.
 XX 04-FEB-2000; 2000US-0180312P.
 XX 26-MAY-2000; 2000US-027456P.
 XX 30-JUN-2000; 2000US-0060808.
 XX 03-AUG-2000; 2000US-00632366.
 XX 21-SEP-2000; 2000US-0234681P.
 XX 27-SEP-2000; 2000US-0236339P.
 XX 04-OCT-2000; 2000GB-00024263.
 XX PA (NOEB-) MOLECULAR DYNAMICS INC.
 XX PI Penn SG, Hanzel DK, Chen W, Rank DR;
 XX DR; 2001-488900/53.
 XX Human genome-derived single exon nucleic acid probes useful for analyzing
 XX gene expression in human bone marrow.
 XX Example 4: SEQ ID NO 30631; 65am - Sequence Listing. English.

SQ Sequence 53 AA;
Query Match 100.0%; Score 29; DB 4; Length 53;
Best Local Similarity 100.0%; Prcd. No. 1.5e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 SSGPSL 6
Db 5 ||||| 10
5 SSGPSL 10

Search completed: March 10, 2004, 09:12:08
Job time : 11.035 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: March 10, 2004, 08:58:54 ; Search time 5.92996 Seconds
(without alignments)
319.245 Million cell updates/sec

Title: US-09-848-834A-6
Perfect Score: 29

Sequence: 1 SSGPSL 6

Scoring table: BIOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SPTREMBL 25;*

- 1: sp_archaea;*
- 2: sp_bacteria;*
- 3: sp_fungi;*
- 4: sp_hexameric;*
- 5: sp_invertebrate;*
- 6: sp_mammal;*
- 7: sp_mhc;*
- 8: sp_organelle;*
- 9: sp_phage;*
- 10: sp_plant;*
- 11: sp_rhodan;*
- 12: sp_virus;*
- 13: sp_vertebrate;*
- 14: sp_inclassified;*
- 15: sp_rvirus;*
- 16: sp_bacteriapl;*
- 17: sp_archeap;*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match	Length	DB ID	Description
1	29	100.0	113	16	Q83F79	Q33F79 coxiella bu
2	29	100.0	174	16	Q8MZ01	Q8mz01 drosophila
3	29	100.0	200	3	Q9UVTS	Q9uvts magnaporthe
4	29	100.0	213	4	Q9UJGL	Q9ujgl homo sapien
5	29	100.0	213	11	Q8VEBL0	Q8vebl0 mus musculu
6	29	100.0	214	11	Q9D8Y9	Q9dy9 mus musculu
7	29	100.0	215	13	Q7SVO6	Q7sy6 xenopus lae
8	29	100.0	226	10	Q9FUJX4	Q9fujx4 arabidopsis
9	29	100.0	257	16	Q8VKB5	Q8vkbs mycobacteri
10	29	100.0	277	4	Q8IXV0	Q8ixv0 homo sapien
11	29	100.0	284	2	Q9L913	Q9l913 aeromonas v
12	29	100.0	348	16	Q8YJL9	Q8yl9 brucella me
13	29	100.0	358	16	Q7U3U5	Q7u3u5 synchococc
14	29	100.0	392	15	Q7XXK1	Q7xxk1 oryza sativ
15	29	100.0	395	4	Q8N4F2	Q8n4f2 homo sapien
16	29	100.0	395	12	Q86606	Q86606 simian virus

RT "Complete genome sequence of the Q-fever pathogen, Coxiella

RT burnetii."

RT "Complete genome sequence of the Q-fever pathogen, Coxiella

RT burnetii."

RT burnetii."</

	Db	79	SSGPSL 84	
RESULT 2 Q8MZQ1 PRELIMINARY; PRT; 174 AA.				
AC Q8MZQ1;	ID Q9UJG1 PRELIMINARY; PRT; 213 AA.			
DT 01-OCT-2002 (TREMBLrel. 22, Created)	ID Q9UJG1;			
DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)	AC Q9UJG1;			
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)	DT 01-MAY-2000 (TREMBLrel. 13, Created)			
DE RE3342P;	DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)			
GN Drosophila melanogaster (Fruit fly).	DB Hypothetical Protein d4_3b4.			
OS Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; OC Phyla; Endopterygota; Diptera; Brachycera; Muscomorpha; OC Ephydriidae; Drosophilidae; Drosophila.	OS Homo sapiens (Human).			
NCBI_TaxID=7227; RN [1]	OC Mammalia; Buteraria; Primates; Catarrhini; Hominidae; Homo. OX NCBI_TaxID=9606;			
RP SEQUENCE FROM N.A.	OX [1]			
RC STRAIN=Berkeley;	RP SEQUENCE FROM N.A.			
RA Stapleton M., Brookstein P., Hong L., Agbayani A., Carlson J., Champé M., Chavez C., Dorsett V., Dressen D., Frise E., George R., Gonzalez M., Guarini H., Krommiller B., Li P., Liao G., Miranda A., Mungall C.J., Munro J., Paragis V., Park S., Patel S., Phoumanenong S., Wan K., Yu C., Lewis S.E., Rubin G.M., Celinker S.;	RA Submitted (JAN-2000) to the EMBL/GenBank/DDBJ databases.			
RL Submitted (MAY-2002) to the EMBL/GenBank/DDBJ databases.	RA TISSUE=Placenta;			
DR EMBL; AY113351; AAC29456.1; -	RA STRAUSBERG R.			
FILEBase; FBgn0063034; BcdNA; RE3342P;	RA Submitted (APR-2001) to the EMBL/GenBank/DDBJ databases.			
SQ 174 AA; 19281 MW; 05C3CE1AB43DB8DA CRC64;	DR EMBL; AL137163; CAB9662.1; -			
Query Match Score 29; DB 5; Length 174; Best Local Similarity 100.0%; Pred. No. 99; Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	DR GO; GO005700.1; -			
Db 122 SSGPSL 6	DR GO; GO005198; F:structural molecule activity; IEA.			
Query Match Score 29; DB 4; Length 213; Best Local Similarity 100.0%; Pred. No. 1.2e+02; Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	DR InterPro; IPR005355; MSP domain.			
Db 122 SSGPSL 127	DR InterPro; IPR008962; PapD-like.			
RESULT 3 Q9UVT8 PRELIMINARY; PRT; 200 AA.	DR PF00635; MSP domain; 1.			
AC Q9UVT8; DT 01-MAY-2000 (TREMBLrel. 13, Created)	DR PROSITE; PS5020.0; MSP; 1.			
DT 01-OCT-2002 (TREMBLrel. 13, Last sequence update)	DR GO; GO005198; F:structural molecule activity; IEA.			
DE 6,7-dimethyl-8-ribityllumazine synthase	DR InterPro; IPR005355; MSP domain.			
OS Magnaporthe grisea (Rice blast fungus) (Pyricularia grisea).	DR InterPro; IPR008962; PapD-like.			
OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes; Sordariomycetes incertae sedis; Magnaporthe; Magnaporthe.	DR RIKEN CDNA 1810018L05 gene.			
NCBI_TaxID=148305; RN [1]	GN 1810018L05RIK.			
RP SEQUENCE FROM N.A.	OS Mus musculus (Mouse).			
RA Person K., Schneidler G., Jordan D.B., Viitanen P.V., Sandalova T.; RT "Comparison of the crystal structures of the pentameric fungal and the icosahedral plant lumazine synthases.";	OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; OC Mammalia; Buteraria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.			
RL Submitted (MAY-1999) to the EMBL/GenBank/DDBJ databases.	OC NCBI_TaxID=10090;			
DR EMBL; AP148449; ADD5372.1; -	OX [1]			
DR PDB; 1C41; 07-AUG-00.	RP SEQUENCE FROM N.A.			
DR GO; GO00092319; C:riboflavin synthase complex; IEA.	RA STRAUSBERG R.			
DR GO; GO0004746; F:riboflavin synthase activity; IEA.	RA Submitted (DEC-2001) to the EMBL/GenBank/DDBJ databases.			
DR GO; GO0009231; P:vitamin B2 biosynthesis; IEA.	DR EMBL; BC018329; AAH18329.1; -			
DR InterPro; IPB002280; DMRL synthase.	DR MGD; MGJ:1911630; 1810018L05Rik.			
DR Pfam; PF00885; DMRL synthase; 1.	DR GO; GO005198; F:structural molecule activity; IEA.			
DR PRODOM; PD003664; DMRL synthase; 1.	DR InterPro; IPR005355; MSP domain.			
DR TIGRFAM; TIGR00014; ribB; 1.	DR InterPro; IPR008962; PapD-like.			
DR SEQUENCE 200 AA; 21072 MW; E079PP49D2227155 CRC64;	DR PROSITE; PS5020.0; MSP; 1.			
Query Match Score 29; DB 3; Length 200; Best Local Similarity 100.0%; Pred. No. 1.1e+02; Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	DR SEQUENCE 213 AA; 24074 MW; FB72756A0528CB34 CRC64;			
Db 1 SSGPSL 6	Query Match Score 29; DB 11; Length 213; Best Local Similarity 100.0%; Pred. No. 1.2e+02; Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;			